



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(21) International Application Number: PCT/US99/18149 (22) International Filing Date: 10 August 1999 (10.08.1999) (30) Priority Data: 60/096,110 10 August 1998 (10.08.1998) US 60/131,352 28 April 1999 (28.04.1999) US (60) Parent Application or Grant NOVIRIO PHARMACEUTICALS LIMITED [/]; (). CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE [/]; (). GOSSELIN, Gilles [/]; (). IMBACH, Jean-Louis [/]; (). BRYANT, Martin, L. [/]; (). GOSSELIN, Gilles [/]; (). IMBACH, Jean-Louis [/]; (). BRYANT, Martin, L. [/]; (). KNOWLES, Sherry; ().		Published
(54) Title: 'beta'-L-2'-DEOXY-NUCLEOSIDES FOR THE TREATMENT OF HEPATITIS B (54) Titre: 'beta'-L-2'-DESOXY-NUCLEOSIDES POUR LE TRAITEMENT DE L'HEPATITE B		
(57) Abstract This invention is directed to a method for treating a host infected with hepatitis B comprising administering an effective amount of an anti-HBV biologically active 2'-deoxy-'beta'-L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof, wherein the 2'-deoxy-'beta'-L-erythro-pentofuranonucleoside has formula (I) wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may be optionally substituted. The 2'-deoxy-'beta'-L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof may be administered either alone or in combination with another 2'-deoxy-'beta'-L-erythro-pentofuranonucleoside or in combination with another anti-hepatitis B agent.		
(57) Abrégé L'invention concerne une méthode de traitement d'un sujet hôte atteint de l'hépatite B consistant à administrer une dose efficace d'un 2'-désoxy-'beta'-L-érythro-pentofuranonucléoside anti-VHB biologiquement actif ou un sel ou un promédicament pharmaceutiquement acceptable de celui-ci, dans laquelle le 2'-désoxy-'beta'-L-érythro-pentofuranonucléoside a la formule (I) dans laquelle R est choisi dans le groupe comprenant H, alkyle à chaîne droite, ramifiée ou cyclique, CO-alkyle, CO-aryle, CO-alkoxyalkyle, CO-aryloxyalkyle, CO-aryle substitué, alkylsulfonyle, arylsulfonyle, aralkylsulfonyle, un reste d'acide aminé, monophosphate, diphosphate ou triphosphate ou un dérivé de phosphate; et BASE représente une base purine ou pyrimidine pouvant être facultativement substituée. Le 2'-désoxy-'beta'-L-érythro-pentofuranonucléoside ou un sel ou un promédicament pharmaceutiquement acceptable de celui-ci peut être administré soit seul soit combiné à un autre 2'-désoxy-'beta'-L-érythro-pentofuranonucléoside ou en combinaison avec un autre agent anti-hépatite B.		

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(54) Title: β -L-2'-DEOXY-NUCLEOSIDES FOR THE TREATMENT OF HEPATITIS B			
<div style="text-align: center;"> <p>(I)</p> </div>			
(57) Abstract This invention is directed to a method for treating a host infected with hepatitis B comprising administering an effective amount of an anti-HBV biologically active 2'-deoxy- β -L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside has formula (I) wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may be optionally substituted. The 2'-deoxy- β -L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof may be administered either alone or in combination with another 2'-deoxy- β -L-erythro-pentofuranonucleoside or in combination with another anti-hepatitis B agent.			

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Description

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β -L-2'-Deoxy-Nucleosides for the Treatment of Hepatitis B

Background of the Invention

This invention is in the area of methods for the treatment of hepatitis B virus (also referred to as "HBV") that includes administering to a host in need thereof, either alone or in combination, an effective amount of one or more of the active compounds disclosed herein, or a pharmaceutically acceptable prodrug or salt of one of these compounds.

HBV is second only to tobacco as a cause of human cancer. The mechanism by which HBV induces cancer is unknown, although it is postulated that it may directly trigger tumor development, or indirectly trigger tumor development through chronic inflammation, cirrhosis, and cell regeneration associated with the infection.

Hepatitis B virus has reached epidemic levels worldwide. After a two to six month incubation period in which the host is unaware of the infection, HBV infection can lead to acute hepatitis and liver damage, that causes abdominal pain, jaundice, and elevated blood levels of certain enzymes. HBV can cause fulminant hepatitis, a rapidly progressive, often fatal form of the disease in which massive sections of the liver are destroyed.

Patients typically recover from acute hepatitis. In some patients, however, high levels of viral antigen persist in the blood for an extended, or indefinite, period, causing a chronic infection. Chronic infections can lead to chronic persistent hepatitis. Patients infected with chronic persistent HBV are most common in developing countries. By mid-1991, there were approximately 225 million chronic carriers of HBV in Asia alone, and worldwide, almost 300 million carriers. Chronic persistent hepatitis can cause fatigue, cirrhosis of the liver, and hepatocellular carcinoma, a primary liver cancer.

In western industrialized countries, high risk groups for HBV infection include those in contact with HBV carriers or their blood samples. The epidemiology of HBV is very similar to that of acquired immune deficiency syndrome (AIDS), which accounts for why HBV infection is common among patients with AIDS or AIDS related complex. However, HBV is more contagious than HIV.

However, more recently, vaccines have also been produced through genetic engineering and are currently used widely. Unfortunately, vaccines cannot help those already infected with HBV. Daily treatments with α -interferon, a genetically engineered protein, has

also shown promise, but this therapy is only successful in about one third of treated patients. Further, interferon cannot be given orally.

A number of synthetic nucleosides have been identified which exhibit activity against HBV. The (-)-enantiomer of BCH-189, known as 3TC, claimed in U. S. Patent 5,539,116 to Liotta, et al., has been approved by the U.S. Food and Drug Administration for the treatment of hepatitis B. See also EPA 0 494 119 A1 filed by BioChem Pharma, Inc.

Cis-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane ("FTC") exhibits activity against HBV. See WO 92/15308; Furman, et al., "The Anti-Hepatitis B Virus Activities, Cytotoxicities, and Anabolic Profiles of the (-) and (+) Enantiomers of cis-5-Fluoro-1-[2-(Hydroxymethyl)-1,3-oxathiolane-5-yl]-Cytosine" Antimicrobial Agents and Chemotherapy, December 1992, page 2686-2692; and Cheng, et al., Journal of Biological Chemistry, Volume 267(20), 13938-13942 (1992).

von Janta-Lipinski et al. disclose the use of the L-enantiomers of 3'-fluoro-modified β -2'-deoxyribonucleoside 5'-triphosphates for the inhibition of hepatitis B polymerases (J. Med. Chem., 1998, 41,2040-2046). Specifically, the 5'-triphosphates of 3'-deoxy-3'-fluoro- β -L-thymidine (β -L-FTTP), 2',3'-dideoxy-3'-fluoro- β -L-cytidine (β -L-FdCTP), and 2',3'-dideoxy-3'-fluoro- β -L-5-methylcytidine (β -L-FMethCTP) were disclosed as effective inhibitors of HBV DNA polymerases.

WO 96/13512 to Genencor International, Inc. and Lipitek, Inc. discloses that certain L-ribofuranosyl nucleosides can be useful for the treatment of cancer and viruses. Specifically disclosed is the use of this class of compounds for the treatment of cancer and HIV.

U. S. Patent Nos. 5,565,438, 5,567,688 and 5,587,362 (Chu, et al.) disclose the use of 2'-fluoro-5-methyl- β -L-arabinofuranolyluridine (L-FMAU) for the treatment of hepatitis B and Epstein Barr virus.

Yale University and University of Georgia Research Foundation, Inc. disclose the use of L-FddC (β -L-5-fluoro-2',3'-dideoxycytidine) for the treatment of hepatitis B virus in WO 92/18517.

The synthetic nucleosides β -L-2'-deoxycytidine (β -L-2'-dC), β -L-2'-deoxythymidine (β -L-dT) and β -L-2'-deoxyadenosine (β -L-2'-dA), are known in the art. Antonin Holy first disclosed β -L-dC and β -L-dT in 1972, "Nucleic Acid Components and Their Analogs. CLIII. Preparation of 2'-deoxy-L-Ribonucleosides of the Pyrimidine Series," *Collect. Czech. Chem.*

5 *Commun.* (1972), 37(12), 4072-87. Morris S. Zedeck et al. first disclosed β -L-dA for the inhibition of the synthesis of induced enzymes in *Pseudomonas testosteroni*, *Mol. Phys.* (1967), 3(4), 386-95.

10 Certain 2'-deoxy- β -L-erythro-pentofuranonucleosides are known to have antineoplastic and selected antiviral activities. Verri et al. disclose the use of 2'-deoxy- β -L-erythro-pentofuranonucleosides as antineoplastic agents and as anti-herpetic agents (*Mol. Pharmacol.* (1997), 51(1), 132-138 and *Biochem. J.* (1997), 328(1), 317-20). Saneyoshi et al. demonstrate the use of 2'-deoxy-L-ribonucleosides as reverse transcriptase (I) inhibitors for the control of retroviruses and for the treatment of AIDS, Jpn. Kokai Tokkyo Koho
15 JP06293645 (1994).

20 Giovanni et al. tested 2'-deoxy- β -L-erythro-pentofuranonucleosides against partially pseudorabies virus (PRV), *Biochem. J.* (1993), 294(2), 381-5.

Chemotherapeutic uses of 2'-deoxy- β -L-erythro-pentofuranonucleosides were studied by Tyrsted et al. (*Biochim. Biophys. Acta* (1968), 155(2), 619-22) and Bloch, et al. (*J. Med. Chem.* (1967), 10(5), 908-12).

25 β -L-2'-deoxythymidine (β -L-dT) is known in the art to inhibit herpes simplex virus type 1 (HSV-1) thymidine kinase (TK). Iotti et al., WO 92/08727, teaches that β -L-dT selectively inhibits the phosphorylation of D-thymidine by HSV-1 TK, but not by human TK. Spaldari et al. reported that L-thymidine is phosphorylated by herpes simplex virus type 1
30 thymidine kinase and inhibits viral growth, *J. Med. Chem.* (1992), 35(22), 4214-20.

In light of the fact that hepatitis B virus has reached epidemic levels worldwide, and has severe and often tragic effects on the infected patient, there remains a strong need to provide new effective pharmaceutical agents to treat humans infected with the virus that have low toxicity to the host.

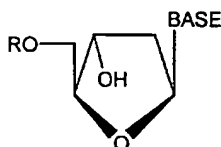
25 Therefore, it is an object of the present invention to provide new methods and compositions for the treatment of human patients or other hosts infected with hepatitis B virus.

Summary of the Invention

A method for the treatment of hepatitis B infection in humans and other host animals is disclosed that includes administering an effective amount of a biologically active 2'-deoxy- β -L-erythro-pentofuranonucleoside (referred to alternatively herein as a β -L-d-nucleoside or a β -L-2'-d-nucleoside) or a pharmaceutically acceptable salt or prodrug thereof, administered either alone or in combination, optionally in a pharmaceutically acceptable carrier. The term 2'-deoxy, as used in this specification, refers to a nucleoside that has no substituent in the 2'-position.

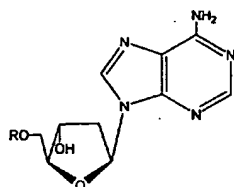
The disclosed 2'-deoxy- β -L-erythro-pentofuranonucleosides, or pharmaceutically acceptable prodrugs or salts or pharmaceutically acceptable formulations containing these compounds are useful in the prevention and treatment of hepatitis B infections and other related conditions such as anti-HBV antibody positive and HBV-positive conditions, chronic liver inflammation caused by HBV, cirrhosis, acute hepatitis, fulminant hepatitis, chronic persistent hepatitis, and fatigue. These compounds or formulations can also be used prophylactically to prevent or retard the progression of clinical illness in individuals who are anti-HBV antibody or HBV-antigen positive or who have been exposed to HBV.

In one embodiment of the present invention, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is a compound of the formula:



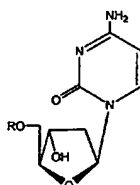
wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may optionally be substituted.

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is β -L-2'-deoxyadenosine or a pharmaceutically acceptable salt or prodrug thereof, of the formula:



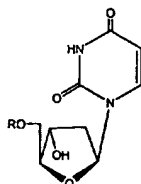
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is β -L-2'-deoxycytidine or pharmaceutically acceptable salt or prodrug thereof of the formula:



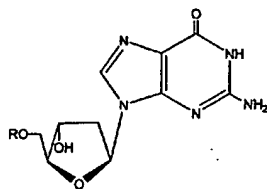
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is β -L-2'-deoxyuridine or pharmaceutically acceptable salt or prodrug thereof of the formula:



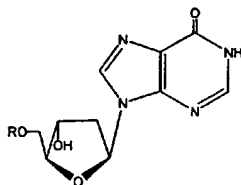
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is β -L-2'-deoxyguanosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



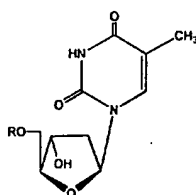
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is β -L-2'-deoxyinosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative is β -L-thymidine or a pharmaceutically acceptable salt or prodrug thereof of the formula:



wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

In another embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside is administered in alternation or combination with one or more other 2'-deoxy- β -L-erythro-

5 pentofuranonucleosides or one or more other compounds which exhibit activity against
hepatitis B virus. In general, during alternation therapy, an effective dosage of each agent is
administered serially, whereas in combination therapy, an effective dosage of two or more
10 agents are administered together. The dosages will depend on absorption, inactivation, and
excretion rates of the drug as well as other factors known to those of skill in the art. It is to
be noted that dosage values will also vary with the severity of the condition to be alleviated.
It is to be further understood that for any particular subject, specific dosage regimens and
15 schedules should be adjusted over time according to the individual need and the professional
judgment of the person administering or supervising the administration of the compositions.

In another embodiment, the invention includes a method for the treatment of humans
infected with HBV that includes administering an HBV treatment amount of a prodrug of the
20 disclosed 2'-deoxy- β -L-erythro-pentofuranonucleoside derivatives. A prodrug, as used
herein, refers to a compound that is converted into the nucleoside on administration in vivo.
Nonlimiting examples include pharmaceutically acceptable salt (alternatively referred to as
"physiologically acceptable salts"), the 5' and N⁴ (cytidine) or N⁶ (adenosine) acylated or
25 alkylated derivatives of the active compound, or the 5'-phospholipid or 5'-ether lipids of the
active compound.

30 Brief Description of the Figures

Figure 1 illustrates a general process for obtaining β -L-erythro-
pentafuranonucleosides (β -L-dN) using L-ribose or L-xylose as a starting material.

35 Figure 2 is a graph which illustrates the metabolism of L-dA, L-dC, and L-dT in
human Hep G2 cells in terms of accumulation and decay. The cells were incubated with 10
 μ M of compound.

40 Figure 3 is a graph which illustrates the antiviral effect of β -L-dA, β -L-dT and β -L-dC
in the woodchuck chronic hepatitis model.

Detailed Description of the Invention

As used herein, the term "substantially in the form of a single isomer" or "in isolated form" refers to a 2'-deoxy- β -L-erythro-pentofuranonucleoside that is at least approximately 95% in the designated stereoconfiguration. In a preferred embodiment, the active compound is administered in at least this level of purity to the host in need of therapy.

As used herein, the term hepatitis B and related conditions refers to hepatitis B and related conditions such as anti-HBV antibody positive and HBV-positive conditions, chronic liver inflammation caused by HBV, cirrhosis, acute hepatitis, fulminant hepatitis, chronic persistent hepatitis, and fatigue. The method of the present invention includes the use of 2'-deoxy- β -L-erythro-pentofuranonucleoside derivatives prophylactically to prevent or retard the progression of clinical illness in individuals who are anti-HBV antibody or HBV-antigen positive or who have been exposed to HBV.

As used herein, the term alkyl, unless otherwise specified, refers to a saturated straight, branched, or cyclic, primary, secondary, or tertiary hydrocarbon, typically of C₁ to C₁₈, preferably C₁ to C₆ and specifically includes but is not limited to methyl, ethyl, propyl, butyl, pentyl, hexyl, isopropyl, isobutyl, sec-butyl, t-butyl, isopentyl, amyl, t-pentyl, cyclopentyl, and cyclohexyl.

As used herein, the term acyl refers to moiety of the formula -C(O)R', wherein R' is alkyl; aryl, alkaryl, aralkyl, heteroaromatic, alkoxyalkyl including methoxymethyl; arylalkyl including benzyl; aryloxyalkyl such as phenoxymethyl; aryl including phenyl optionally substituted with halogen, C₁ to C₄ alkyl or C₁ to C₄ alkoxy, or the residue of an amino acid. The term acyl specifically includes but is not limited to acetyl, propionyl, butyryl, pentanoyl, 3-methylbutyryl, hydrogen succinate, 3-chlorobenzoate, benzoyl, acetyl, pivaloyl, mesylate, propionyl, valeryl, caproic, caprylic, capric, lauric, myristic, palmitic, stearic, and oleic.

As used herein, the term purine or pyrimidine base, includes, but is not limited to, 6-alkylpurine and N⁶-alkylpurines, N⁶-acylpurines, N⁶-benzylpurine, 6-halopurine, N⁶-vinylpurine, N⁶-acetylenic purine, N⁶-acyl purine, N⁶-hydroxyalkyl purine, N⁶-thioalkyl purine, N²-alkylpurines, N⁴-alkylpyrimidines, N⁴-acylpyrimidines, 4-benzylpyrimidine, N⁴-halopyrimidines, N⁴-acetylenic pyrimidines, 4-acyl and N⁴-acyl pyrimidines, 4-hydroxyalkyl pyrimidines, 4-thioalkyl pyrimidines, thymine, cytosine, 6-azapyrimidine, including 6-azacytosine, 2- and/or 4-mercaptopyrimidine, uracil, C⁵-alkylpyrimidines, C⁵-

5 benzylpyrimidines, C⁵-halopyrimidines, C⁵-vinylpyrimidine, C⁵-acetylenic pyrimidine, C⁵-
acyl pyrimidine, C⁵-hydroxyalkyl purine, C⁵-amidopyrimidine, C⁵-cyanopyrimidine, C⁵-
10 nitropyrimidine, C⁵-aminopyrimidine, N²-alkylpurines, N²-alkyl-6-thiopurines, 5-
azacytidinyl, 5-azauracilyl, triazolopyridinyl, imidazolopyridinyl, pyrrolopyrimidinyl, and
pyrazolopyrimidinyl. Functional oxygen and nitrogen groups on the base can be protected as
15 necessary or desired. Suitable protecting groups are well known to those skilled in the art,
and include trimethylsilyl, dimethylhexylsilyl, *t*-butyldimethylsilyl, and *t*-butyldiphenylsilyl,
trityl, alkyl groups, acyl groups such as acetyl and propionyl, methanesulfonyl, and *p*-
toluenesulfonyl.

The term biologically active nucleoside, as used herein, refers to a nucleoside which
20 exhibits an EC₅₀ of 15 micromolar or less when tested in 2.2.15 cells transfected with the
hepatitis virion.

Preferred bases include cytosine, 5-fluorocytosine, 5-bromocytosine, 5-iodocytosine,
uracil, 5-fluorouracil, 5-bromouracil, 5-iodouracil, 5-methyluracil, thymine, adenine,
25 guanine, inosine, xanthine, 2,6-diaminopurine, 6-aminopurine, 6-chloropurine and 2,6-
dichloropurine, 6-bromopurine, 2,6-dibromopurine, 6-iodopurine, 2,6-di-iodopurine, 5-
bromovinylcytosine, 5-bromovinyluracil, 5-bromoethenylcytosine, 5-bromoethenyluracil, 5-
trifluoromethylcytosine, 5-trifluoromethyluracil.

The 2'-deoxy-β-L-erythro-pentofuranonucleoside can be provided as a 5'
30 phospholipid or a 5'-ether lipid, as disclosed in the following references: Kucera, L.S., N.
Lyer, E. Leake, A. Raben, Modest E.J., D. L.W., and C. Piantadosi. 1990. Novel
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Marasco C.J., S.L. morris-Natschke, K.L. Meyer, F. Gumus, J.R. Surles, K.S. Ishaq, L.S.
Kucera, N. lyer, C.A. Wallen, S. Piantadosi, and E.J. Modest. 1991-Synthesis and evaluation
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40 34:1408-1414; Hostetler, K.Y., D.D. Richman, D.A. Carson, L.M. Stuhmiller, G.M. T. van
Wijk, and H. van den Bosch. 1992. Greatly enhanced inhibition of human immunodeficiency
virus type 1 replication in CEM and HT4-6C cells by 31-deoxythymidine diphosphate
45 dimyristoylglycerol, a lipid prodrug of 31-deoxythymidine. Antimicrob Agents Chemother.
36:2025-2029; Hostetler, K.Y., L.M. Stuhmiller, H.B. Lenting, H. van den Bosch, and D.D.

5 Richman. 1990. Synthesis and antiretroviral activity of phospholipid analogs of
azidothymidine and other antiviral nucleosides. J. Biol Chem. 265:6112-7.

10 The 2'-deoxy- β -L-erythro-pentofuranonucleoside can be converted into a
pharmaceutically acceptable ester by reaction with an appropriate esterifying agent, for
example, an acid halide or anhydride. The nucleoside or its pharmaceutically acceptable
prodrug can be converted into a pharmaceutically acceptable salt thereof in a conventional
manner, for example, by treatment with an appropriate base or acid. The ester or salt can be
converted into the parent nucleoside, for example, by hydrolysis.

15 As used herein, the term pharmaceutically acceptable salts or complexes refers to salts
or complexes of the 2'-deoxy- β -L-erythro-pentofuranonucleosides that retain the desired
biological activity of the parent compound and exhibit minimal, if any, undesired
toxicological effects. Nonlimiting examples of such salts are (a) acid addition salts formed
20 with inorganic acids (for example, hydrochloric acid, hydrobromic acid, sulfuric acid,
phosphoric acid, nitric acid, and the like), and salts formed with organic acids such as acetic
acid, oxalic acid, tartaric acid, succinic acid, malic acid, ascorbic acid, benzoic acid, tannic
acid, palmoic acid, alginic acid, polyglutamic acid, naphthalenesulfonic acids,
25 naphthalenedisulfonic acids, and polygalacturonic acid; (b) base addition salts formed with
cations such as sodium, potassium, zinc, calcium, bismuth, barium, magnesium, aluminum,
copper, cobalt, nickel, cadmium, sodium, potassium, and the like, or with an organic cation
formed from N,N-dibenzylethylene-diamine, ammonium, or ethylenediamine; or (c)
30 combinations of (a) and (b); e.g., a zinc tannate salt or the like.

35 The term prodrug, as used herein, refers to a compound that is converted into the
nucleoside on administration in vivo. Nonlimiting examples are pharmaceutically acceptable
salts (alternatively referred to as "physiologically acceptable salts"), the 5' and N⁴ or N⁶
acylated or alkylated derivatives of the active compound, and the 5'-phospholipid and 5'-
ether lipid derivatives of the active compound.

40 Modifications of the active compounds, specifically at the N⁴, N⁶ and 5'-O positions,
can affect the bioavailability and rate of metabolism of the active species, thus providing
control over the delivery of the active species.

45 A preferred embodiment of the present invention is a method for the treatment of
HBV infections in humans or other host animals, that includes administering an effective
amount of one or more of a 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative selected

5 from the group consisting of β -L-2'-deoxyadenosine, β -L-2'-deoxycytidine, β -L-2'-
deoxyuridine, β -L-2'-guanosine, β -L-2'-deoxyinosine, and β -L-2'-deoxythymidine, or a
physiologically acceptable prodrug thereof, including a phosphate, 5' and or N⁶ alkylated or
acylated derivative, or a physiologically acceptable salt thereof, optionally in a
10 pharmaceutically acceptable carrier. The compounds of this invention either possess
anti-HBV activity, or are metabolized to a compound or compounds that exhibit anti-HBV
activity. In a preferred embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside is
administered substantially in the form of a single isomer, i.e., at least approximately 95% in
15 the designated stereoconfiguration.

Nucleotide Prodrugs

20 Any of the nucleosides described herein can be administered as a stabilized nucleotide
prodrug to increase the activity, bioavailability, stability or otherwise alter the properties of
the nucleoside. A number of nucleotide prodrug ligands are known. In general, alkylation,
acylation or other lipophilic modification of the mono, di or triphosphate of the nucleoside
25 will increase the stability of the nucleotide. Examples of substituent groups that can replace
one or more hydrogens on the phosphate moiety are alkyl, aryl, steroids, carbohydrates,
including sugars, 1,2-diacylglycerol and alcohols. Many are described in R. Jones and N.
Bischofberger, *Antiviral Research*, 27 (1995) 1-17. Any of these can be used in combination
30 with the disclosed nucleosides to achieve a desired effect.

In one embodiment, the 2'-deoxy- β -L-erythro-pentofuranonucleoside is provided as
5'-hydroxyl lipophilic prodrug. Nonlimiting examples of U.S. patents that disclose suitable
35 lipophilic substituents that can be covalently incorporated into the nucleoside, preferably at
the 5'-OH position of the nucleoside or lipophilic preparations, include U.S. Patent Nos.
5,149,794 (Sep. 22, 1992, Yatvin et al.); 5,194,654 (Mar. 16, 1993, Hostetler et al.,
5,223,263 (June 29, 1993, Hostetler et al.); 5,256,641 (Oct. 26, 1993, Yatvin et al.);
40 5,411,947 (May 2, 1995, Hostetler et al.); 5,463,092 (Oct. 31, 1995, Hostetler et al.);
5,543,389 (Aug. 6, 1996, Yatvin et al.); 5,543,390 (Aug. 6, 1996, Yatvin et al.); 5,543,391
(Aug. 6, 1996, Yatvin et al.); and 5,554,728 (Sep. 10, 1996; Basava et al.).

45 Foreign patent applications that disclose lipophilic substituents that can be attached to
the 2'-deoxy- β -L-erythro-pentofuranonucleoside derivative of the present invention, or
lipophilic preparations, include WO 89/02733, WO 90/00555, WO 91/16920, WO 91/18914,

5 WO 93/00910, WO 94/26273, WO 96/15132, EP 0 350 287, EP 93917054.4, and WO
91/19721.

Additional nonlimiting examples of 2'-deoxy- β -L-erythro-pentofuranonucleosides are
those that contain substituents as described in the following publications. These derivatized
10 2'-deoxy- β -L-erythro-pentofuranonucleosides can be used for the indications described in the
text or otherwise as antiviral agents, including as anti-HBV agents. Ho, D.H.W. (1973)
Distribution of kinase and deaminase of 1 β -D-arabinofuranosylcytosine in tissues of man
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enzymatic synthesis of 5'-(3-sn-phosphatidyl)nucleosides and their antileukemic activities.
Chem. Pharm. Bull. 36, 209-217. One preferred phosphate prodrug group is the S-acyl-2-
20 thioethyl group, also referred to as "SATE."

Combination or Alternation Therapy

25 It has been recognized that drug-resistant variants of HBV can emerge after prolonged
treatment with an antiviral agent. Drug resistance most typically occurs by mutation of a
gene that encodes for an enzyme used in the viral life cycle, and most typically in the case of
HBV, DNA polymerase. Recently, it has been demonstrated that the efficacy of a drug
30 against HBV infection can be prolonged, augmented, or restored by administering the
compound in combination or alternation with a second, and perhaps third, antiviral compound
that induces a different mutation from that caused by the principle drug. Alternatively, the
pharmacokinetics, biodistribution, or other parameter of the drug can be altered by such
35 combination or alternation therapy. In general, combination therapy is typically preferred
over alternation therapy because it induces multiple simultaneous stresses on the virus.

40 The anti-hepatitis B viral activity of β -L-2'-dA, β -L-2'-dC, β -L-2'-dU, β -L-2'-dG, β -
L-2'-dT, β -L-dI, or other β -L-2'-nucleosides provided herein, or the prodrugs, phosphates, or
salts of these compounds, can be enhanced by administering two or more of these nucleosides
in combination or alternation. Alternatively, for example, one or more of β -L-2'-dA, β -L-2'-
45 dC, β -L-2'-dU, β -L-2'-dG, β -L-2'-dT, β -L-dI, or other β -L-2'-nucleosides provided herein
can be administered in combination or alternation with 3TC, FTC, L-FMAU, DAPD,
famciclovir, penciclovir, BMS-200475, bis pom PMEA (adefovir, dipivoxil); lobucavir,
ganciclovir, or ribavarin.

5 In any of the embodiments described herein, if the β -L-2'-nucleoside of the present invention is administered in combination or alternation with a second nucleoside or nonnucleoside reverse transcriptase inhibitor that is phosphorylated to an active form, it is preferred that a second compound be phosphorylated by an enzyme that is different from that
10 which phosphorylates the selected β -L-2'-nucleoside of the present invention in vivo. Examples of kinase enzymes are thymidine kinase, cytosine kinase, guanosine kinase, adenosine kinase, deoxycytidine kinase, 5'-nucleotidase, and deoxyguanosine kinase.

15 Preparation of the Active Compounds

The 2'-deoxy- β -L-erythro-pentofuranonucleoside derivatives of the present invention are known in the art and can be prepared according to the method disclosed by Holy, *Collect. Czech. Chem. Commun.* (1972), 37(12), 4072-87 and *Mol. Phys.* (1967), 3(4), 386-95.

20 A general process for obtaining β -L-erythro-pentafuranonucleosides (β -L-dN) is shown in Figure 1, using L-ribose or L-xylose as a starting material.

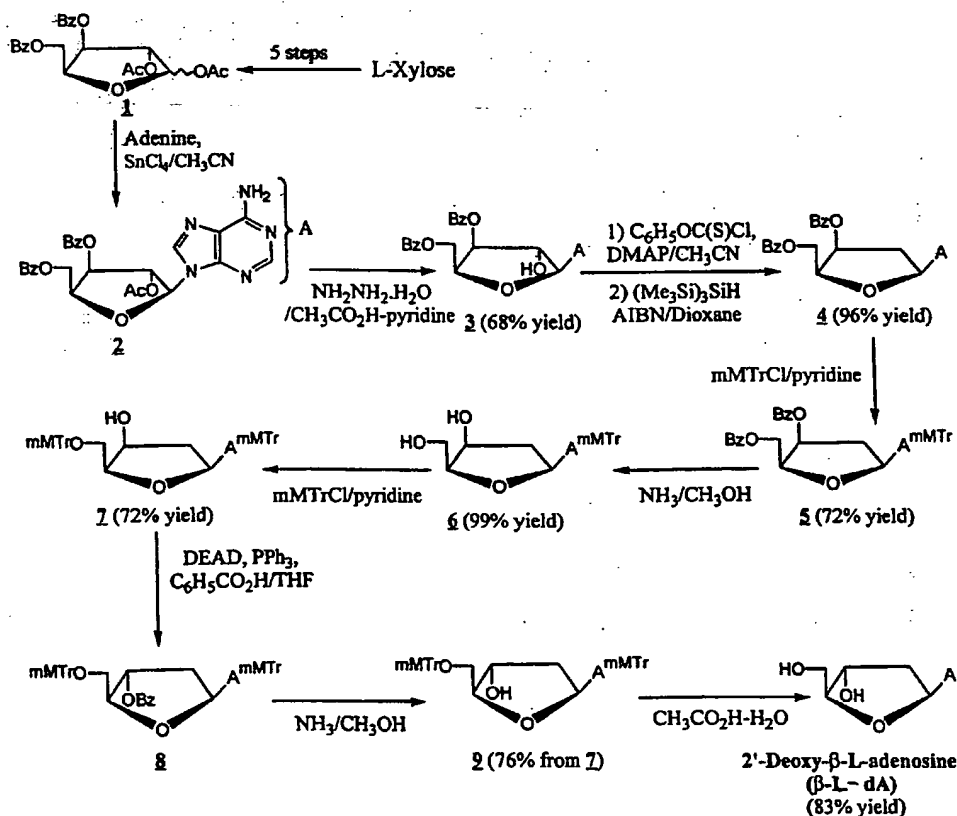
25 Mono, di, and triphosphate derivatives of the active nucleosides can be prepared as described according to published methods. The monophosphate can be prepared according to the procedure of Imai et al., *J. Org. Chem.*, 34(6), 1547-1550 (June 1969). The diphosphate can be prepared according to the procedure of Davisson et al., *J. Org. Chem.*, 52(9), 1794-1801 (1987). The triphosphate can be prepared according to the procedure of Hoard et al., *J. Am. Chem. Soc.*, 87(8), 1785-1788 (1965).

30 Experimental Protocols

Melting points were determined in open capillary tubes on a Gallenkamp MFB-595-010 M apparatus and are uncorrected. The UV absorption spectra were recorded on an
35 Uvikon 931 (KONTRON) spectrophotometer in ethanol. $^1\text{H-NMR}$ spectra were run at room temperature in $\text{DMSO-}d_6$ with a Bruker AC 250 or 400 spectrometer. Chemical shifts are given in ppm, $\text{DMSO-}d_5$ being set at 2.49 ppm as reference. Deuterium exchange, decoupling
40 experiments or 2D-COSY were performed in order to confirm proton assignments. Signal multiplicities are represented by s (singlet), d (doublet), dd (doublet of doublets), t (triplet), q (quadruplet), br (broad), m (multiplet). All J-values are in Hz. FAB mass spectra were
45 recorded in the positive- ($\text{FAB} > 0$) or negative- ($\text{FAB} < 0$) ion mode on a JEOL DX 300 mass spectrometer. The matrix was 3-nitrobenzyl alcohol (NBA) or a mixture (50:50, v/v) of glycerol and thioglycerol (GT). Specific rotations were measured on a Perkin-Elmer 241

spectropolarimeter (path length 1 cm) and are given in units of $10^{-1} \text{ deg cm}^2 \text{ g}^{-1}$. Elemental analysis were carried out by the "Service de Microanalyses du CNRS, Division de Vernaison" (France). Analyses indicated by the symbols of the elements or functions were within $\pm 0.4\%$ of theoretical values. Thin layer chromatography was performed on precoated aluminium sheets of Silica Gel 60 F₂₅₄ (Merck, Art. 5554), visualisation of products being accomplished by UV absorbency followed by charring with 10% ethanolic sulfuric acid and heating. Column chromatography was carried out on Silica Gel 60 (Merck, Art. 9385) at atmospheric pressure.

Example 1 Stereospecific Synthesis of 2'-Deoxy- β -L-Adenosine



9-(3,5-Di-O-benzoyl- β -L-xylofuranosyl)adenine (3)

A solution of 9-(2-O-acetyl-3,5-di-O-benzoyl- β -L-xylofuranosyl)adenine **2** [Ref.: Gosselin, G.; Bergogne, M.-C.; Imbach, J.-L., "Synthesis and Antiviral Evaluation of β -L-Xylofuranosyl Nucleosides of the Five Naturally Occurring Nucleic Acid Bases", *Journal of Heterocyclic Chemistry*, 1993, **30** (Oct.-Nov.), 1229-1233] (8.30 g, 16.05 mmol) and hydrazine hydrate 98% (234 mL, 48.5 mmol) in a mixture of pyridine / glacial acetic acid (4/1, v/v, 170 mL) was stirred at room temperature for 22 h. The reaction was quenched by adding acetone (40 mL) and stirring was continued for one additional hour. The reaction mixture was reduced to one half of its volume, diluted with water (250 mL) and extracted with chloroform (2 x 150 mL). The organic layer was washed successively with an aqueous saturated solution of NaHCO₃ (3 x 100 mL) and water (3 x 100 mL), dried, filtered, concentrated and co-evaporated with toluene and methanol. The residue was purified by silica gel column chromatography (0-3% MeOH in dichloromethane) to give **3** (5.2 g, 68%) precipitated from diisopropyl ether : ¹H NMR (DMSO-*d*₆) : δ 4.5-4.9 (m, 4H, H-2', H-4', H-5' and H-5''), 5.64 (t, 1H, H-3', J_{2',3'} = J_{3',4'} = 3.5 Hz), 6.3 (br s, 1H, OH-2'), 6.45 (d, 1H, H-1', J_{1',2'} = 4.6 Hz), 7.3 (br s, 2H, NH₂-6), 7.4-7.9 (m, 10H, 2 benzoyls), 8.07 and 8.34 (2s, 2H, H-2 and H-8); ms : matrix G/T, (FAB⁺) m/z 476 [M+H]⁺, 136 [BH₂]⁺, (FAB⁻) m/z 474 [M-H]⁻, 134 [B]⁻; UV (95% ethanol) : λ_{max} 257 nm (ϵ 16400), 230 nm (ϵ 29300), λ_{min} 246 nm (ϵ 14800); [α]_D²⁰ = - 64 (c 1.07, CHCl₃). Anal. Calcd for C₂₄H₂₁N₅O₄ (M = 475.45) : C, 60.43; H, 4.45; N, 14.73. Found : C, 60.41; H, 4.68; N, 14.27.

9-(3,5-Di-O-benzoyl-2-deoxy- β -L-threo-pentofuranosyl)adenine (4).

To a solution of compound **3** (1.00 g, 2.11 mmol) in dry acetonitrile (65 mL) were added 4-(dimethylamino)pyridine (0.77 g, 6.32 mmol) and phenoxythiocarbonyl chloride (0.44 mL, 3.16 mmol). The mixture was stirred at room temperature for 2 h. After concentration, the residue was dissolved in dichloromethane (50 mL) and washed successively with water (2 x 30 mL), aqueous solution of hydrochloric acid 0.5 N (30 mL) and water (3 x 30 mL). The organic layer was dried, filtered and concentrated to dryness. The crude thiocarbonylated intermediate was directly treated with tris-(trimethylsilyl)silane hydride (0.78 mL, 5.23 mmol) and α,α' -azoisobutyronitrile (AIBN, 0.112 g, 0.69 mmol) in dry dioxane (17 mL) at reflux for 2 h. The solvent was removed under vacuum and the residue was purified by silica gel column chromatography (0-5% MeOH in dichloromethane) to give pure **4** (0.93 g, 96%) as a foam : ¹H NMR (DMSO-*d*₆) : δ 2.9-3.1 (m, 2H, H-2' and H-

2"), 4.6 - 4.7 (m, 3H, H-4', H-5' and H-5"), 5.8 (br s, 1H, H-3'), 6.43 (dd, 1H, H-1', $J_{1',2'} = 3.1$ Hz, $J_{1',2''} = 7.6$ Hz), 7.3 (br s, 2H, NH₂-6), 7.4-7.9 (m, 10H, 2 benzoyls), 8.05 and 8.33 (2s, 2H, H-2 and H-8); ms : matrix G/T, (FAB⁺) m/z 460 [M+H]⁺, 325 [S]⁺, 136 [BH₂]⁺, (FAB⁻) m/z 458 [M-H]⁻, 134 [B]⁻; UV (95% ethanol) : λ_{\max} 261 nm (ϵ 14400), 231 nm (ϵ 26300), λ_{\min} 249 nm (ϵ 12000); $[\alpha]_D^{20} = -38$ (c 1.04, DMSO).

6-*N*-(4-Monomethoxytrityl)-9-(3,5-di-*O*-benzoyl-2-deoxy- β -L-threo-pentofuranosyl)adenine (5).

To a solution of compound **4** (0.88 g, 1.92 mmol) in dry pyridine (40 mL) was added 4-monomethoxytrityl chloride (1.18 g, 3.84 mmol). The mixture was stirred at 60°C for 24 h. After addition of methanol (5 mL), the solution was concentrated to dryness, the residue was dissolved in dichloromethane (50 mL) and washed successively with water (30 mL), aqueous saturated NaHCO₃ (30 mL) and water (30 mL). The organic layer was dried, filtered, concentrated and co-evaporated with toluene to give pure **5** (1.01 g, 72%) as a foam : ¹H NMR (CDCl₃) : δ 2.9-3.0 (m, 2H, H-2' and H-2"), 3.62 (s, 3H, OCH₃), 4.6-4.8 (m, 3H, H-4', H-5' and H-5"), 5.85 (pt, 1H, H-3'), 6.44 (dd, 1H, H-1', $J_{1',2'} = 3.1$ Hz, $J_{1',2''} = 7.3$ Hz), 6.9 (br s, 1H, NH-6), 6.7-6.8 and 7.2-7.4 (2m, 24H, 2 benzoyls and MMTr), 7.97 and 8.13 (2s, 2H, H-2 and H-8); ms : matrix G/T, (FAB⁺) m/z 732 [M+H]⁺, (FAB⁻) m/z 730 [M-H]⁻; UV (95% ethanol) : λ_{\max} 274 nm (ϵ 12100), 225 nm (ϵ 24200), λ_{\min} 250 nm (ϵ 5900); $[\alpha]_D^{20} = -16$ (c 1.12, DMSO).

6-*N*-(4-Monomethoxytrityl)-9-(2-deoxy- β -L-threo-pentofuranosyl)-adenine (6).

Compound **5** (0.95 g, 1.30 mmol) was treated with a solution (saturated at -10°C) of methanolic ammonia (40 mL), at room temperature overnight. After concentration, the residue was dissolved in dichloromethane (60 mL) and washed with water (30 mL). The aqueous layer was extracted twice with dichloromethane (10 mL). The combined organic layer was dried, filtered and concentrated. The residue was purified by silica gel column chromatography (0-5% MeOH in dichloromethane) to give pure **6** (0.67 g, 98%) as a foam : ¹H NMR (CDCl₃) : δ 2.6-2.9 (m, 2H, H-2' and H-2"), 3.5 (br s, 1H, OH-5'), 3.55 (s, 3H, OCH₃), 3.9-4.0 (m, 3H, H-4', H-5' and H-5"), 4.5-4.6 (m, 1H, H-3'), 6.03 (dd, 1H, H-1', $J_{1',2'} = 4.0$ Hz, $J_{1',2''} = 8.8$ Hz), 7.0 (br s, 1H, NH-6), 6.7-6.8 and 7.1-7.4 (2m, 14H, MMTr), 7.40 (d, 1H, OH-3', $J_{H,OH} = 10.6$ Hz), 7.80 and 7.99 (2s, 2H, H-2 and H-8); ms : matrix G/T, (FAB⁺) m/z 524 [M+H]⁺, 408 [BH₂]⁺, (FAB⁻) m/z 1045 [2M-H]⁻, 522 [M-H]⁻, 406 [B]⁻; UV (95% ethanol) : λ_{\max} 275 nm (ϵ 12300), λ_{\min} 247 nm (ϵ 3600); $[\alpha]_D^{20} = +28$ (c 0.94, DMSO).

6-*N*-(4-Monomethoxytrityl)-9-(2-deoxy-5-*O*-(4-monomethoxytrityl)- β -L-threo-pentofuranosyl)adenine (7).

Compound **6** (0.62 g, 1.24 mmol) in dry pyridine (25 mL) was treated with 4-monomethoxytrityl chloride (0.46 g, 1.49 mmol) at room temperature for 16 h. After addition of methanol (5 mL), the mixture was concentrated to dryness. The residue was dissolved in dichloromethane (60 mL) and washed successively with water (40 mL), a saturated aqueous solution of NaHCO₃ (40 mL) and water (3 x 40 mL). The organic layer was dried, filtered, concentrated and co-evaporated with toluene and methanol. The residue was purified by silica gel column chromatography (0-10% MeOH in dichloromethane) to give **7** (0.71 g, 72%) as a foam: ¹H NMR (DMSO-*d*₆): δ 2.21 (d, 1H, H-2' J_{2',2''} = 14.3 Hz), 2.6-2.7 (m, 1H, H-2''), 3.1-3.3 (2m, 2H, H-5' and H-5''), 3.64 and 3.65 (2s, 6H, 2 x OCH₃), 4.1-4.2 (m, 1H, H-4'), 4.2-4.3 (m, 1H, H-3'), 5.68 (d, 1H, OH-3', J_{H,OH} = 5.2 Hz), 6.24 (d, 1H, H-1', J_{1',2'} = 7.0 Hz), 6.7-6.8 and 7.1-7.3 (2m, 29H, 2 MMTTr and NH-6), 7.83 and 8.21 (2s, 2H, H-2 and H-8); ms: matrix G/T, (FAB⁺) m/z 796 [M+H]⁺, 408 [BH₂]⁺, (FAB⁻) m/z 794 [M-H]⁻, 406 [B]⁻; UV (95% ethanol): λ_{max} 275 nm (ϵ 30900), λ_{min} 246 nm (ϵ 12800); [α]_D²⁰ = +14 (c 1.03, DMSO).

6-*N*-(4-Monomethoxytrityl)-9-(3-*O*-benzoyl-2-deoxy-5-*O*-(4-monomethoxytrityl)- β -L-erythro-pentofuranosyl)adenine (8).

A solution of diethylazodicarboxylate (0.38 mL, 2.49 mmol) in dry tetrahydrofuran (20 mL) was added dropwise to a cooled solution (0°C) of nucleoside **7** (0.66 g, 0.83 mmol), triphenylphosphine (0.66 g, 2.49 mmol) and benzoic acid (0.30 g, 2.49 mmol) in dry THF (20 mL). The mixture was stirred at room temperature for 18 h and methanol (1 mL) was added. The solvents were removed under reduced pressure and the crude material was purified by silica gel column chromatography (0-5% ethyl acetate in dichloromethane) to give compound **8** slightly contaminated by triphenylphosphine oxide.

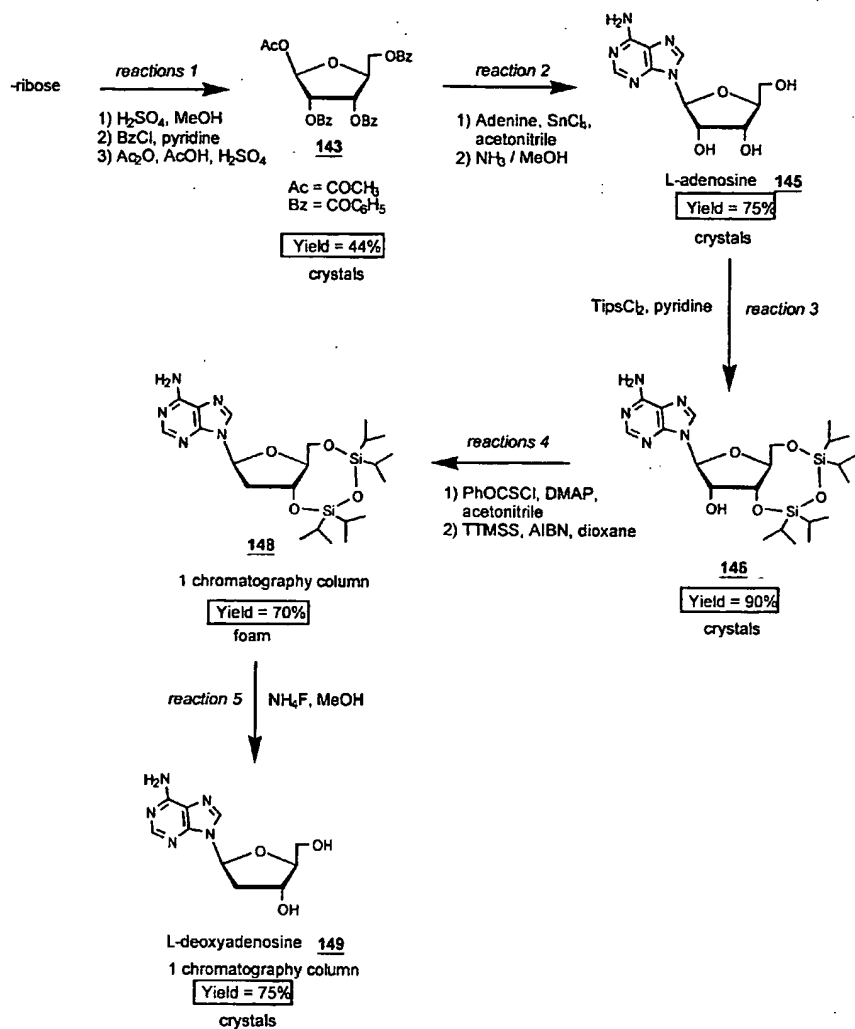
6-*N*-(4-Monomethoxytrityl)-9-(2-deoxy-5-*O*-(4-monomethoxytrityl)- β -L-erythro-pentofuranosyl)adenine (9).

Compound **8** was treated by a solution (saturated at -10°C) of methanolic ammonia (20 mL), at room temperature for 24 h, then the reaction mixture was concentrated to dryness. The residue was dissolved in dichloromethane (30 mL) and washed with water (20 mL). The aqueous layer was extracted by dichloromethane (2 x 20 mL) and the combined organic phase was dried, filtered and concentrated. Pure compound **9** (0.50 g, 76% from **7**) was

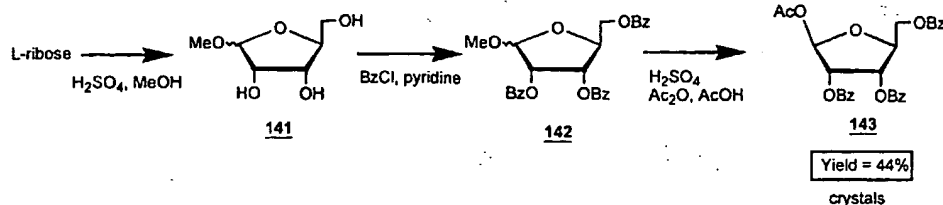
obtained as a foam after purification by silica gel column chromatography (0-2% MeOH in dichloromethane): ^1H NMR ($\text{DMSO}-d_6$): δ 2.2-2.3 (m, 1H, H-2'), 2.8-2.9 (m, 1H, H-2''), 3.1-3.2 (m, 2H, H-5' and H-5''), 3.64 and 3.65 (2s, 6H, 2 x OCH_3), 3.97 (pq, 1H, H-4'), 4.4-4.5 (m, 1H, H-3'), 5.36 (d, 1H, OH-3', $J_{\text{H,OH}} = 4.5$ Hz), 6.34 (t, 1H, H-1', $J_{1',2'} = J_{1',2''} = 6.4$ Hz), 6.8-6.9 and 7.1-7.4 (2m, 29H, 2 MMTr and NH-6), 7.81 and 8.32 (2s, 2H, H-2 and H-8); ms: matrix G/T, (FAB $^+$) m/z 796 $[\text{M}+\text{H}]^+$, 408 $[\text{BH}_2]^+$, (FAB $^-$) m/z 794 $[\text{M}-\text{H}]^-$, 406 $[\text{B}]^-$; UV (95% ethanol): λ_{max} 276 nm (ϵ 42600), λ_{min} 248 nm (ϵ 23300); $[\alpha]_{\text{D}}^{20} = +29$ (c 1.05, DMSO).

2'-Deoxy- β -L-adenosine (β -L-dA)

Compound **9** (0.44 g, 0.56 mmol) was treated with an aqueous solution of acetic acid 80% (17 mL) at room temperature for 5 h. The mixture was concentrated to dryness, the residue was dissolved in water (20 mL) and washed with diethyl ether (2 x 15 mL). The aqueous layer was concentrated and co-evaporated with toluene and methanol. The desired 2'-deoxy- β -L-adenosine (β -L-dA) (0.12 g, 83%) was obtained after purification by silica gel column chromatography (0-12% MeOH in dichloromethane) and filtration through a Millex HV-4 unit (0.45 μ , Millipore): mp 193-194°C (crystallized from water) (Lit. 184-185°C for L-enantiomer [Ref.: Robins, M. J.; Khwaja, T. A.; Robins, R. K. *J. Org. Chem.* **1970**, *35*, 636-639] and 187-189°C for D-enantiomer [Ref.: Ness, R. K. in *Synthetic Procedures in Nucleic Acid Chemistry*; Zorbach, W.W., Tipson, R. S., Eds.; J. Wiley and sons: New York, **1968**; Vol 1, pp 183-187]; ^1H NMR ($\text{DMSO}-d_6$): δ 2.2-2.3 and 2.6-2.7 (2m, 2H, H-2' and H-2''), 3.4-3.6 (2m, 2H, H-5' and H-5''), 3.86 (pq, 1H, H-4'), 4.3-4.4 (m, 1H, H-3'), 5.24 (t, 1H, OH-5', $J_{\text{H,OH}} = 5.8$ Hz), 5.30 (d, 1H, OH-3', $J_{\text{H,OH}} = 4.0$ Hz), 6.32 (dd, 1H, H-1', $J_{1',2'} = 6.2$ Hz, $J_{1',2''} = 7.8$ Hz), 7.3 (br s, 2H, NH-6), 8.11 and 8.32 (2s, 2H, H-2 and H-8); ms: matrix G/T, (FAB $^+$) m/z 252 $[\text{M}+\text{H}]^+$, 136 $[\text{BH}_2]^+$, (FAB $^-$) m/z 250 $[\text{M}-\text{H}]^-$, 134 $[\text{B}]^-$; UV (95% ethanol): λ_{max} 258 nm (ϵ 14300), λ_{min} 226 nm (ϵ 2100); $[\alpha]_{\text{D}}^{20} = +25$ (c 1.03, H_2O), (Lit. $[\alpha]_{\text{D}}^{20} = +23$ (c 1.0, H_2O) for L-enantiomer [Ref.: Robins, M. J.; Khwaja, T. A.; Robins, R. K. *J. Org. Chem.* **1970**, *35*, 636-639] and $[\alpha]_{\text{D}}^{20} = -25$ (c 0.47, H_2O) for D-enantiomer [Ref.: Ness, R. K. in *Synthetic Procedures in Nucleic Acid Chemistry*; Zorbach, W.W., Tipson, R. S., Eds.; J. Wiley and sons: New York, **1968**; Vol 1, pp 183-187]). Anal. Calcd for $\text{C}_{10}\text{H}_{13}\text{N}_5\text{O}_3 + 1.5 \text{H}_2\text{O}$ ($M = 278.28$): C, 43.16; H, 5.80; N, 25.17. Found: C, 43.63; H, 5.45; N, 25.33.

Example 2 Stereoselective Synthesis of 2'-Deoxy- β -L-Adenosine (β -L-dA)

Reaction 1:



Precursor: L-ribose (Cultor Science Food, CAS [24259-59-4], batch RIB9711013)

Reactants: Sulphuric acid 95-97% (Merck; ref 1.00731.1000); Benzoyl chloride (Fluka; ref 12930); Sodium sulfate (Prolabo; ref 28111.365)

Solvents: Methanol P.A. (Prolabo; ref 20847.295); Pyridine 99% (Acros; ref 131780025); Dichloromethane P.A. (Merck; ref 1.06050.6025); Acetic acid P.A. (carlo erba; ref 20104298); Acetic anhydride (Fluka; ref 45830); Ethanol 95 (Prolabo; ref 20823.293)

References: Recondo, E. F., and Rinderknecht, H., Eine neue, Einfache Synthese des 1-O-Acetyl-2,3,5-Tri-O-β-D-Ribofuranosides. *Helv. Chim. Acta*, 1171-1173 (1959).

A solution of L-ribose **140** (150 g, 1 mol) in methanol (2 liters) was treated with sulphuric acid (12 ml) and left at +4°C for 12hrs, and then neutralised with pyridine (180 ml). Evaporation gave an α,β mixture of methyl ribofuranosides **141** as a syrup. A solution of this anomeric mixture in pyridine (1.3 liters) was treated with benzoyl chloride (580 ml, 5 mol) with cooling and mechanical stirring. The solution was left at room temperature for 12 hrs and then poured on ice (about 10 liters) with continued stirring. The mixture (an oil in water) was filtered on a Cellite bed. The resulting oil on the cellite bed was washed with water (3x3 liters) and then dissolved with ethyl acetate (3 liters). The organic phase was washed with a 5% NaHCO₃ solution (2 liters) and water (2 liters), dried over sodium sulfate, filtered and evaporated to give 1-O-methyl-2,3,5-tri-O-benzoyl-α/β-L-ribofuranose **142** as a thick syrup. The oil was dissolved in acetic anhydride (560 ml) and acetic acid (240 ml). The solution was, after the dropwise addition of concentrated sulphuric acid (80 ml), kept in the cold (+4°C) under mechanical stirring for 10 hrs. The solution was then poured on ice (about 10 liters) under continued stirring. The mixture (oily compound in water) was filtered on a Cellite bed. The resulting gummy solid on the cellite bed was washed with water (3x3 liters) and then dissolved in dichloromethane (2,5 liters). The organic phase was washed with 5% NaHCO₃ (1 liter) and water (2x2 liters), dried over sodium sulfate, filtered and evaporated to

give a gummy solid **143**, which was crystallized from ethanol 95 (yield 225 g, 44%).

Analyses for 1-O-acetyl-2,3,5-tri-O-benzoyl-β-L-ribofuranose **143:**

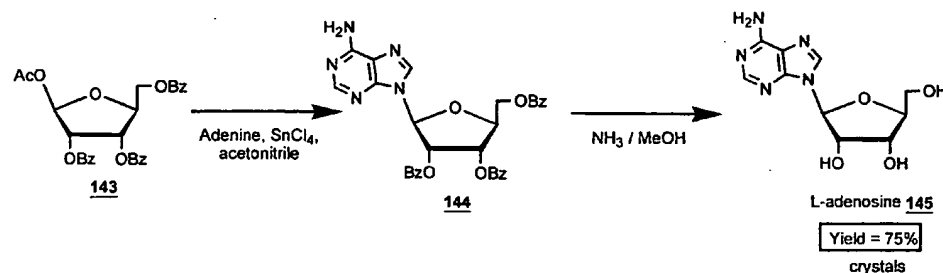
mp 129-130°C (EtOH 95) (lit.(1) mp 130-131°C)

¹H NMR (200 MHz, CDCl₃): δ 8.09-7.87 (m, 6H, H_{Arom}), 7.62-7.31 (m, 9H, H_{Arom}), 6.43 (s, 1H, H₁), 5.91 (dd, 1H, H₃, J_{3,4} 6.7 Hz; J_{3,2} 4.9 Hz), 5.79 (pd, 1H, H₂, J_{2,3} 4.9 Hz; J_{1,2} <1), 4.78 (m, 2H, H₄ and H₅), 4.51 (dd, 1H, H₅, J_{5,5'} 13.1 Hz, J_{5',4} 5.5 Hz), 2.00 (s, 3H, CH₃CO); (identical to commercial 1-O-acetyl-2,3,5-tri-O-benzoyl-β-D-ribofuranose)

Mass analysis (FAB+, GT) m/z 445 (M-OAc)+

Elemental analysis C₂₈H₂₄O₉ Calculated C 66.66 H 4.79; found C H

Reaction 2:



Precursor: Adenine (Pharma-Waldhof; ref 400134.001 lot 45276800)

Reactants: Stannic chloride fuming (Fluka; ref 96558); NH₃ / Methanol (methanol saturated with NH₃; see page 5); Sodium sulfate (Prolabo; ref 28111.365)

Solvents: Acetonitrile (Riedel-de Hean; ref 33019; distilled over CaH₂); Chloroform Pur (Acros; ref 22706463); Ethyl acetate Pur (Carlo erba; ref 528299)

References: Saneyoshi, M., and Satoh, E., Synthetic Nucleosides and Nucleotides. XIII. Stannic Chloride Catalyzed Ribosylation of Several 6-Substituted Purines. *Chem. Pharm. Bull.*, **27**, 2518-2521 (1979).; Nakayama, C., and Saneyoshi, M., Synthetic Nucleosides and Nucleotides. XX. Synthesis of Various 1-β-Xylofuranosyl-5-Alkyluracils and Related Nucleosides. *Nucleosides, Nucleotides*, **1**, 139-146 (1982).

Adenine (19.6 g, 144 mmol) was suspended in acetonitrile (400 ml) with 1-O-acetyl-2,3,5-tri-O-benzoyl-β-L-ribofuranose **143** (60 g, 119 mmol). To this suspension was added stannic chloride fuming (22 ml, 187 mmol). After 12 hrs, the reaction was concentrated to a

small volume (about 100 ml), and sodium hydrogencarbonate (110 g) and water (120 ml) were added. The resulting white solid (tin salts) was extracted with hot chloroform (5x200 ml). The combined extracts were filtered on a cellite bed. The organic phase was washed with a NaHCO₃ 5% solution and water, dried over sodium sulfate, filtered and evaporated to give compound **144** (60 g, colorless foam). The foam was treated with methanol saturated with ammonia (220 ml) in sealed vessel at room temperature under stirring for 4 days. The solvent was evaporated off under reduced pressure and the resulting powder was suspended in ethyl acetate (400 ml) at reflux for 1 hr. After filtration, the powder was recrystallized from water (220 ml) to give L-adenosine **145** (24 g, crystals, 75%)

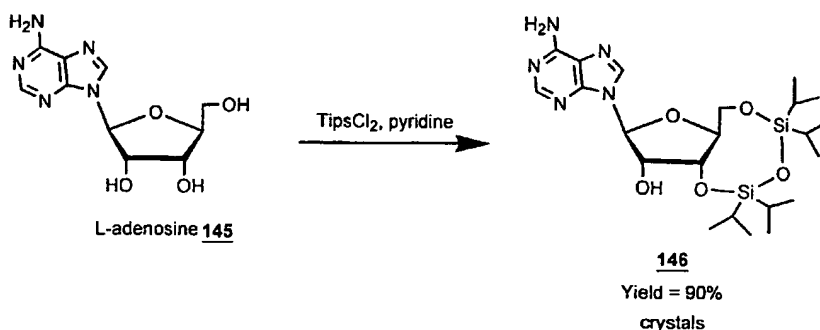
Analyses for β -L-adenosine:

mp 233-234°C (water) (lit.(4) mp 235°-238°C)

¹H NMR (200 MHz, DMSO-D₆): δ 8.34 and 8.12 (2s, 2H, H₂ and H₈), 7.37 (1s, 2H, NH₂), 5.86 (d, 1H, H_{1'}, J_{1'2'} 6.2 Hz), 5.43 (m, 2H, OH_{2'} and OH_{5'}), 5.19 (d, 1H, OH_{3'}, J 3.7 Hz), 4.60 (m, H_{2'}), 4.13 (m, 1H, H_{3'}), 3.94 (m, 1H, H_{4'}), 3.69-3.49 (m, 2H, H_{5'a} and H_{5'b}), (identical to commercial D-adenosine)

Mass analysis (FAB+, GT) m/z 268 (M+H)⁺, 136(BH₂)⁺

Reaction 3:



Reactants: 1,3-Dichloro-1,1,3,3-tetraisopropylidisiloxane (Fluka; ref 36520); Sodium sulfate (Prolabo; ref 28111.365)

Solvents: Pyridine 99% (Acros; ref 131780025); Ethyl acetate Pur (Carlo erba; ref 528299); Acetonitrile (Riedel-de Haen; ref 33019)

Reference : Robins, M.J., et al., Nucleic Acid Related Compounds. 42. A General Procedure for the Efficient Deoxygenation of Secondary Alcohols. Regiospecific and

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"TTMSS" (Fluka; ref 93411); α,α' -Azobisisobutyronitrile "AIBN" (Fluka, ref 11630); Sodium sulfate (Prolabo; ref 28111.365)

Solvents: Acetonitrile (Riedel-de Haen; ref 33019); Ethyl acetate Pur (Carlo Erba; ref 528299); Dioxan P.A. (Merck; ref 1.09671.1000); Dichloromethane (Merck; ref 1.06050.6025); Methanol (Carlo Erba; ref 309002);

Reference: Robins, M. J., Wilson, J. S., and Hansske, F., Nucleic Acid Related Compounds. 42. A General Procedure for the Efficient Deoxygenation of Secondary Alcohols. Regiospecific and Stereoselective Conversion of Ribonucleosides to 2'-Deoxynucleosides. *J. Am. Chem. Soc.*, **105**, 4059-4065 (1983).

To compound **146** (34 g, 67 mmol) were added acetonitrile (280 ml), DMAP (16.5 g, 135 mmol) and phenyl chlorothionocarbonate (10.2 ml, 73 mmol). The solution was stirred at room temperature for 12 hrs. Solvent was evaporated and the residue was partitioned between ethyl acetate (400 ml) and a HCl 0.5N solution (400 ml). The organic layer was washed with a HCl 0.5N solution (400 ml) and water (2x400 ml), dried over sodium sulfate, filtered and evaporated to dryness to give the intermediate as a pale yellow solid. The crude **147** was dissolved in dioxan (ml) and AIBN (3.3 g, 20 mmol) and TTMSS (33 ml, 107 mmol) were added. The solution was progressively heated until reflux and stirred for 2 hrs. The reaction was concentrated to a yellow oil which was chromatographed (eluent dichloromethane/methanol 95/5) to give compound **148** (23 g, colorless foam, 70%). An aliquot was crystallized from ethanol/ petroleum ether.

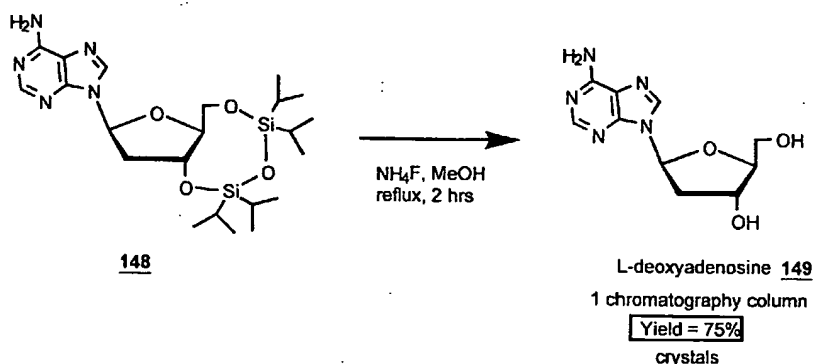
Analyses for 3',5'-O-(1,1,3,3-tetraisopropyl-1,3-disiloxanyl)-2'-deoxy- β -L-adenosine **148:**

mp 110-111°C (EtOH/petroleum ether) (Lit.(5) mp 113-114°C (EtOH))

^1H NMR (200 MHz, CDCl_3): δ 8.33 and 8.03 (2s, 2H, H_2 and H_8), 6.30 (dd, 1H, H_1 , J 2.85 Hz, J 7.06 Hz), 5.63 (sl, 2H, NH_2), 4.96 (m, 1H, H_3), 4.50 (m, 2H, $\text{H}_{5'a}$ and $\text{H}_{5'b}$), 2.68 (m, 2H, $\text{H}_{2'a}$ and $\text{H}_{2'b}$), 1.08 (m, 28H, isopropyl protons)

Mass analysis (FAB+, GT) m/z 494 ($\text{M}+\text{H}$) $^+$, 136 (BH_2) $^+$

Reaction 5:



Reactants : Ammonium fluoride (Fluka; ref 09742); Silica gel (Merck; ref 1.07734.2500)

Solvents : Methanol P.A. (Prolabo; ref 20847.295); Dichloromethane P.A. (Merck; ref 1.06050.6025); Ethanol 95 (Prolabo; ref 20823.293)

Reference : Zhang, W., and Robins, M. J., Removal of Silyl Protecting Groups from Hydroxyl Functions with Ammonium Fluoride in Methanol. *Tetrahedron Lett.*, **33**, 1177-1180 (192).

A solution of 3',5'-O-(1,1,3,3-tetraisopropyl-1,3-disiloxanyl)-2'-deoxy-L-adenosine **148** (32 g, 65 mmol) and ammonium fluoride (32 g, mmol) in methanol was stirred at reflux for 2 hrs. Silica gel was added and the mixture was carefully evaporated to give a white powder. This powder was added on the top of a silica column, which was eluted with dichloromethane/methanol 9/1. The appropriate fractions were combined and evaporated to give a white powder, which was crystallized from ethanol 95 (12.1 g, 75%).

Analyses for 2'-Deoxy- β -L-adenosine 149 :

mp 189-190°C (EtOH 95) (identical to commercial 2'-deoxy-D-adenosine)

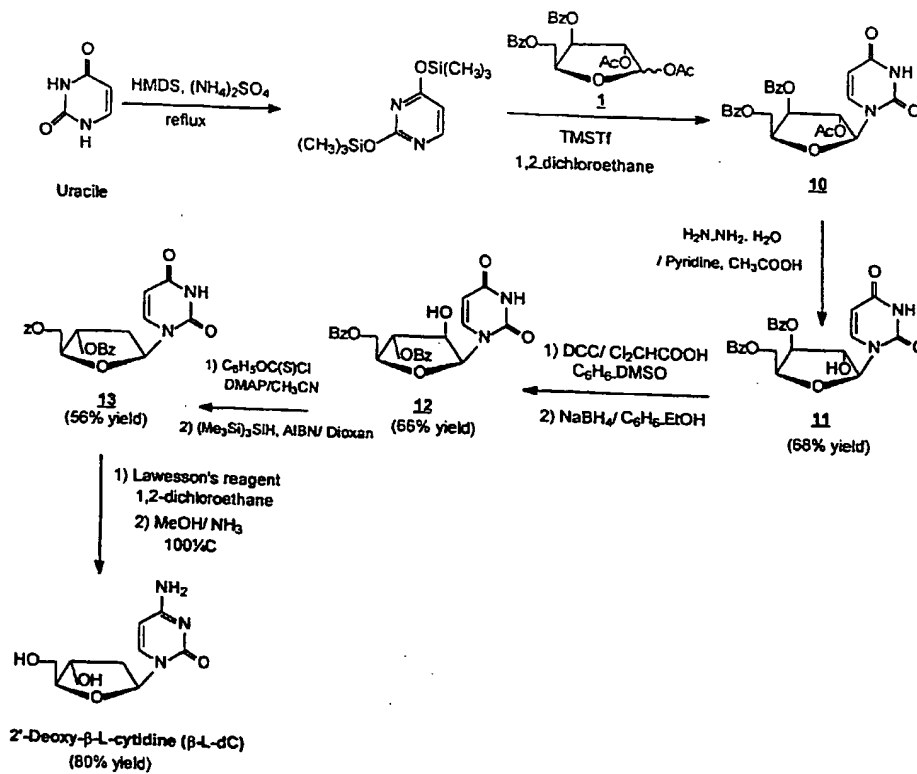
^1H NMR (200 MHz, DMSO- D_6): δ 8.35 and 8.14 (2s, 2H, H_2 and H_8), 7.34 (sl, 2H, NH_2), 6.35 (dd, 1H, H_1 , J 6.1 Hz, J 7.85 Hz), 5.33 (d, 1H, OH_2 , J 4.0 Hz), 5.28 (dd, 1H, H_3 , J 4.95 Hz; J 6.6 Hz), 4.42 (m, 1H, OH_5'), 3.88 (m, 1H, H_4), 3.63-3.52 (m, 2H, $\text{H}_{5'a}$ and $\text{H}_{5'b}$), 2.71 (m, 1H, $\text{H}_{2'a}$), 2.28 (m, 1H, $\text{H}_{2'b}$). (identical to commercial 2'-deoxy-D-adenosine)

$\alpha_D +26^\circ$ (c 0.5 water) (commercial 2'-deoxy-D-adenosine -25° (c 0.5 water)).

UV λ_{max} 260 nm (ϵ 14100) (H_2O).

Mass analysis (FAB+, GT) m/z 252 ($\text{M}+\text{H}$) $^+$, 136 (BH_2) $^+$

Example 3 Stereospecific Synthesis of 2'-Deoxy-β-L-Cytidine



1-(3,5-Di-O-benzoyl- β -L-xylofuranosyl)uracil (11)

Hydrazine hydrate (1.4 mL, 28.7 mmol) was added to a solution of 1-(2-O-acetyl-3,5-di-O-benzoyl- β -L-xylofuranosyl)uracil **10** [Ref.: Gosselin, G.; Bergogne, M.-C.; Imbach, J.-L., "Synthesis and Antiviral Evaluation of β -L-Xylofuranosyl Nucleosides of the Five Naturally Occurring Nucleic Acid Bases", Journal of Heterocyclic Chemistry, 1993, 30 (Oct.-Nov.), 1229-1233] (4.79 g, 9.68 mmol) in pyridine (60 mL) and acetic acid (15 mL). The solution was stirred overnight at room temperature. Acetone was added (35 mL) and the mixture was stirred for 30 min. The reaction mixture was evaporated under reduced pressure. The resulting residue was purified by silica gel column chromatography [eluent: stepwise gradient of methanol (0-4%) in dichloromethane to give **11** (3.0 g, 68%) which was crystallized from cyclohexane/dichloromethane: mp = 111-114°C; $^1\text{H-NMR}$ (DMSO- d_6): δ 11.35 (br s, 1H, NH), 7.9-7.4 (m, 11H, 2 C₆H₅CO, H-6), 6.38 (d, 1H, OH-2', J_{OH-2'} = 4.2 Hz), 5.77 (d, 1H, H-1', J_{1'-2'} = 1.9 Hz), 5.55 (d, 1H, H-5, J₅₋₆ = 8 Hz), 5.54 (dd, 1H, H-3', J_{3'-2'} = 3.9 Hz and J_{3'-4'} = 1.8 Hz), 4.8 (m, 1H, H-4'), 4.7 (m, 2H, H-5' and H-5''), 4.3 (m, 1H, H-2''); MS: FAB>0 (matrix GT) m/z 453 (M+H)⁺, 105 (C₆H₅CO)⁺; FAB<0 (matrix GT) m/z 451 (M-H)⁻, 121 (C₆H₅CO₂)⁻, 111 (B); Anal. Calcd for C₂₃H₂₀N₂O₈·H₂O : C, 58.09 ; H, 4.76 ; N, 5.96. Found : C, 57.71 ; H, 4.42 ; N, 5.70.

1-(3,5-Di-O-benzoyl- β -L-arabinofuranosyl)uracil (12)

To a solution of 1-(3,5-di-O-benzoyl- β -L-xylofuranosyl)uracil **11** (8 g, 17.7 mL) in an anhydrous benzene-DMSO mixture (265 mL, 6:4, v/v) were added anhydrous pyridine (1.4 mL), dicyclohexylcarbodiimide (10.9 g, 53 mmol) and dichloroacetic acid (0.75 mL). The resulting mixture was stirred at room temperature for 4h, then diluted with ethyl acetate (400 mL) and a solution of oxalic acid (4.8 g, 53 mmol) in methanol (14 mL) was added. After being stirred for 1h, the solution was filtered. The filtrate was washed with a saturated NaCl solution (2x500mL), 3% NaHCO₃ solution (2x500mL) and water (2x500mL). The organic phase was dried over Na₂SO₄, then evaporated under reduced pressure. The resulting residue was then solubilized in an EtOH absolute-benzene mixture (140 mL, 2 : 1, v/v). To this solution at 0°C was added NaBH₄ (0.96 g, 26.5 mmol). After being stirred for 1h, the solution was diluted with ethyl acetate (400 mL), then filtered. The filtrate was washed with a saturated NaCl solution (400 mL) and water (400 mL). The organic phase was dried over Na₂SO₄, then evaporated under reduced pressure. The resulting crude material was purified by silica gel column chromatography [eluent: stepwise gradient of methanol (0-3%) in

dichloromethane to give **12** (5.3 g, 66%) which was crystallized from acetonitrile: mp = 182-183°C; ¹H-NMR (DMSO-*d*₆): δ 11.35 (br s, 1H, NH), 8.0-7.5 (m, 11H, 2 C₆H₅CO, H-6), 6.23 (br s, 1H, OH-2'), 6.15 (d, 1H, H-1', J_{1,2'} = 4 Hz), 5.54 (d, 1H, H-5, J_{5,6} = 8.1 Hz), 5.37 (t, 1H, H-3', J_{3,2'} = J_{3,4'} = 2.6 Hz), 4.7-4.6 (m, 2H, H-5' and H-5''), 4.5 (m, 1H, H-4'), 4.4 (m, 1H, H-2'); MS: FAB>0 (matrix GT) m/z 453 (M+H)⁺, 341 (S)⁺, 113 (BH₂)⁺, 105 (C₆H₅CO)⁺; FAB<0 (matrix GT) m/z 451 (M-H)⁻, 121 (C₆H₅CO₂)⁻, 111 (B)⁻; Anal. Calcd for C₂₃H₂₀N₂O₈: C, 61.06; H, 4.46; N, 6.19. Found: C, 60.83; H, 4.34; N, 6.25.

1-(3,5-Di-O-benzoyl-2-deoxy-β-L-erythro-pentofuranosyl)uracil (13**)**

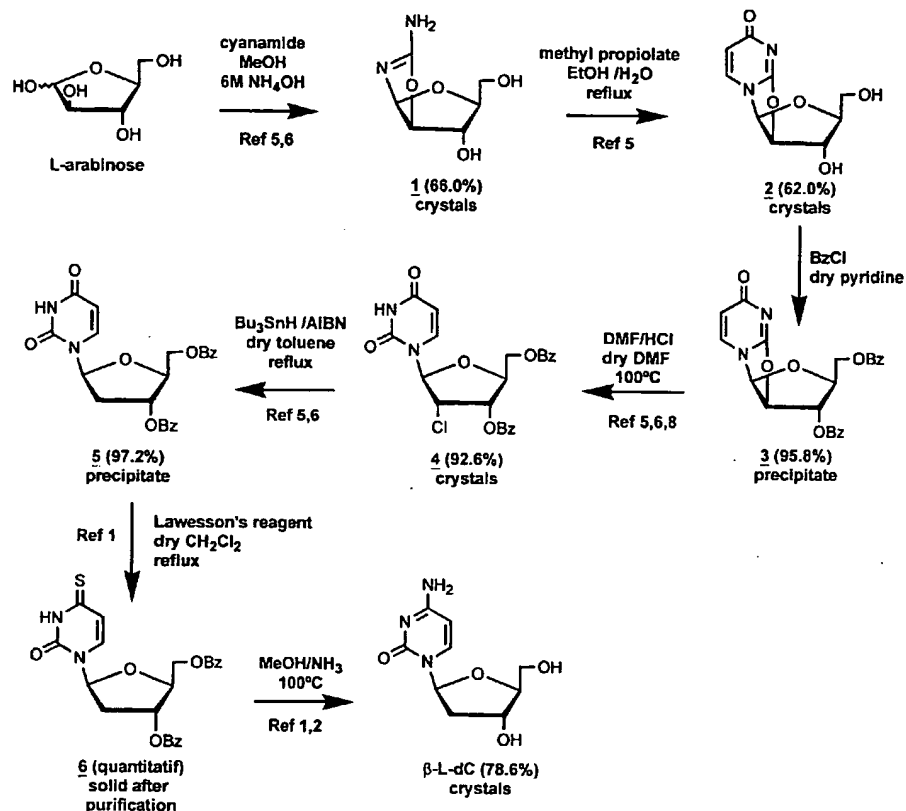
To a solution of 1-(3,5-di-O-benzoyl-β-L-arabinofuranosyl)uracil **12** (5.2 g, 11.4 mmol) in anhydrous 1,2-dichloroethane (120 mL) were added phenoxythiocarbonyl chloride (4.7 mL, 34.3 mL) and 4-(dimethylamino)pyridine (DMAP, 12.5 g, 102.6 mmol). The resulting solution was stirred at room temperature under argon atmosphere for 1h and then evaporated under reduced pressure. The residue was dissolved in dichloromethane (300 mL) and the organic solution was successively washed with an ice-cold 0.2 N hydrochloric acid solution (3x 200 mL) and water (2x200 mL), dried over Na₂SO₄ then evaporated under reduced pressure. The crude material was co-evaporated several times with anhydrous dioxane and dissolved in this solvent (110 mL). To the resulting solution were added under argon tris-(trimethylsilyl)silane hydride (4.2 mL, 13.7 mmol) and α,α'-azoisobutyronitrile (AIBN, 0.6 g, 3.76 mmol). The reaction mixture was heated and stirred at 100°C for 1h under argon, then cooled to room temperature and evaporated under reduced pressure. The residue was purified by silica gel column chromatography [eluent: stepwise gradient of methanol (0-5%)] to give **13** (2.78 g, 56%) which was crystallized from EtOH: mp = 223-225°C; ¹H-NMR (DMSO-*d*₆): δ 11.4 (br s, 1H, NH), 8.0-7.5 (m, 11H, 2 C₆H₅CO, H-6), 6.28 (t, 1H, H-1', J = 7 Hz), 5.5 (m, 2H, H-1' and H-5), 4.6-4.4 (m, 3H, H-4', H-5' and H-5''), 2.6 (m, 2H, H-2' and H-2''); MS: FAB>0 (matrix GT) m/z 437 (M+H)⁺, 3325 (S)⁺; FAB<0 (matrix GT) m/z 435 (M-H)⁻, 111 (B)⁻; Anal. Calcd for C₂₃H₂₀N₂O₇: C, 63.30; H, 4.62; N, 6.42. Found: C, 62.98; H, 4.79; N, 6.40.

2'-Deoxy-β-L-cytidine (β-L-dC)

Lawesson's reagent (1.72 g, 4.26 mmol) was added under argon to a solution of 1-(3,5-di-O-benzoyl-2-deoxy-β-L-erythro-pentofuranosyl)uracil **13** (2.66 g, 6.1 mmol) in anhydrous 1,2-dichloroethane (120mL) and the reaction mixture was stirred under reflux for 2h. The solvent was then evaporated under reduced pressure and the residue was purified by

5 silica gel column chromatography [eluent: stepwise gradient of ethyl acetate (0-8%) in
dichloromethane] to give the 4-thio intermediate as a yellow foam. A solution of this thio-
intermediate (1.5 g, 3.31 mmol) in methanolic ammonia (previously saturated at -10°C and
10 tightly stopped) (50 mL) was heated at 100°C in a stainless-steel bomb for 3h and then cooled
to 0°C. The solution was evaporated under reduced pressure. The resulting crude material
was purified by silica gel column chromatography [eluent: stepwise gradient of methanol(0-
20%) in dichloromethane]. Finally, the appropriate fractions were pooled, filtered through a
unit Millex HV-4 (0.45 µm, Millipore) and evaporated under reduced pressure to provide the
15 desired 2'-deoxy-β-L-cytidine (β-L-dC) as a foam (0.6 g, 80%) which was crystallized from
absolute EtOH: mp=198-199°C; ¹H-NMR (DMSO-*d*₆): δ 7.77 (d, 1H, H-6, *J*_{6,5} = 7.4 Hz),
7.10 (br d, 2H, NH₂), 6.13 (t, 1H, H-1', *J* = 6.7 Hz), 5.69 (d, 1H, H-5, *J*_{5,6} = 7.4 Hz), 5.19 (d,
20 1H, OH-3', *J*_{OH-3'} = 4.1 Hz), 4.96 (t, 1H, OH-5', *J*_{OH-5'} = *J*_{OH-5''} = 5.2 Hz), 4.1 (m, 1H, H-3'),
3.75 (m, 1H, H-4'), 3.5 (m, 2H, H-5' and H-5''), 2.0 (m, 1H, H-2'), 1.9 (m, 1H, H-2''); MS:
FAB>0 (matrix GT) *m/z* 228 (M+H)⁺, 112 (BH₂)⁺; FAB<0 (matrix GT) *m/z* 226(M-H)⁻;
[α]²⁰_D = - 69 (c 0.52, DMSO) [[α]²⁰_D = + 76 (c 0.55, DMSO) for a commercially available
25 hydrochloride salt of the D-enantiomer]. Anal. Calcd for C₉H₁₃N₃O₄: C, 47.57 ; H, 5.77 ; N,
18.49. Found : C, 47.35 ; H, 5.68 ; N, 18.29.

Example 4 Stereoselective Synthesis of 2'-Deoxy-β-L-Cytidine (β-L-dC)



2-Amino-β-L-arabinofurano[1',2':4,5]oxazoline (**1**)

A mixture of L-arabinose (170 g, 1.13 mol), cyanamide (100g, 2.38 mol), methanol (300 ml), and 6M-NH₄OH (50 ml) was stirred at room temperature for 3 days and then kept at -10°C overnight. The product was collected with suction, washed successively with methanol and ether, and dried *in vacuo*. Yield, 130 g (66.0%) of the analytically pure compound **1**, m.p. 170-172°C; ¹H NMR (DMSO-*d*₆) δ ppm 6.35 (br s, 2H, NH₂), 5.15 (d, 1H, H-1, J = 5.6 Hz), 5.45 (br s, 1H, OH-3), 4.70 (br s, 1H, OH-5), 4.55 (d, 1H, H-2, J = 5.6 Hz), 4.00 (br s, 1H, H-3), 3.65 (m, 1H, H-4), 3.25 (m, 2H, H-5, H-5').

Reagents:**L-arabinose:** Fluka, >99.5%, ref 10839**Cyanamide:** Fluka, >98%, ref 28330***O*^{2,2'}-anhydro-β-L-uridine (2)**

A solution of compound **1** (98.8 g, 0.57 mol) and methyl propiolate (98 ml) in 50% aqueous ethanol (740 ml) was refluxed for 5h, then cooled and concentrated under diminished pressure to the half of the original volume. After precipitation with acetone (600 ml), the product was collected with suction, washed with ethanol and ether, and dried. The mother liquor was partially concentrated, the concentrate precipitated with acetone (1000 ml), the solid collected with suction, and washed with acetone and ether to afford another crop of the product. Over-all yield, 80 g (62%) of compound **2**, m.p. 236-240°C; ¹H NMR (DMSO-*d*₆) δ ppm 7.87 (d, 1H, *H*-6, *J* = 7.4 Hz), 6.35 (d, 1H, *H*-1', *J* = 5.7 Hz), 5.95 (d, 1H, *H*-5, *J* = 7.4 Hz), 5.90 (d, 1H, *OH*-3'), 5.20 (d, 1H, *H*-2', *J* = 5.7 Hz), 5.00 (m, 1H, *OH*-3'), 4.44 (br s, 1H, *H*-3'), 4.05 (m, 1H, *H*-4'), 3.25 (m, 2H, *H*-5, *H*-5').

Reagent:**Methyl propiolate:** Fluka, >97%, ref 81863**3',5'-Di-*O*-benzoyl-*O*^{2,2'}-anhydro-β-L-uridine (3)**

To a solution of compound **2** (71.1 g, 0.31 mol) in anhydrous pyridine (1200 ml) was added benzoyl chloride (80.4 ml) at 0°C and under argon. The reaction mixture was stirred at room temperature for 5 h under exclusion of atmospheric moisture and stopped by addition of ethanol. The solvents were evaporated under reduced pressure and the resulting residue was coevaporated with toluene and absolute ethanol. The crude mixture was then diluted with ethanol and the precipitate collected with suction, washed successively with ethanol and ether, and dried. Yield, 129 g (95.8%) of compound **3**, m.p. 254 °C; ¹H NMR (DMSO-*d*₆) δ ppm 8.1-7.4 (m, 11H, C₆H₅CO, *H*-6), 6.50 (d, 1H, *H*-1', *J* = 5.7 Hz), 5.90 (d, 1H, *H*-5, *J* = 7.5 Hz), 5.80 (d, 1H, *H*-2', *J* = 5.8 Hz), 5.70 (d, 1H, *H*-3') 4.90 (m, 1H, *H*-4'), 4.35 (m, 2H, *H*-5, *H*-5').

Reagent:**Benzoyl chloride:** Fluka, p.a., ref 12930

3',5'-Di-O-benzoyl-2'-chloro-2'-deoxy- β ,L-uridine (4)

To a solution of compound **3** (60.3 g, 0.139 mol) in dimethylformamide (460 ml) was added at 0°C a 3.2 N-HCl/DMF solution (208 ml, prepared *in situ* by adding 47.2 ml of acetyl chloride at 0°C to a solution of 27.3 ml of methanol and 133.5 ml of dimethylformamide). The reaction mixture was stirred at 100°C for 1h under exclusion of atmospheric moisture, cooled down, and poured into water (4000 ml). The precipitate of compound **4** was collected with suction, washed with water, and recrystallised from ethanol. The crystals were collected, washed with cold ethanol and ether, and dried under diminished pressure. Yield, 60.6 g (92.6%) of compound **4**, m.p. 164-165°C; ¹H NMR (DMSO-*d*₆) δ ppm 8.7 (br s, 1H, NH), 8.1-7.3 (m, 11H, C₆H₅CO, *H*-6), 6.15 (d, 1H, *H*-1', *J* = 4.8 Hz), 5.5 (m, 2H, *H*-5, *H*-2'), 4.65 (m, 4H, *H*-3', *H*-4', *H*-5', *H*-5'').

Reagent:

Acetyl chloride: Fluka, p.a., ref 00990

3',5'-Di-O-benzoyl-2'-deoxy- β ,L-uridine (5)

A mixture of compound **4** (60.28 g, 0.128 mol), tri-*n*-butyltin hydride (95 ml) and azabisisobutyronitrile (0.568 g) in dry toluene (720 ml) was refluxed under stirring for 5 h and cooled down. The solid was collected with suction and washed with cold toluene and petroleum ether. The filtrate was concentrated under reduced pressure and diluted with petroleum ether to deposit an additional crop of compound **5**. Yield, 54.28 g (97.2%) of compound **5**; m.p. 220-221°C; ¹H NMR (CDCl₃) δ ppm 8.91 (br s, 1H, NH), 8.1-7.5 (m, 11H, C₆H₅CO and *H*-6), 6.43 (q, 1H, *H*-1', *J*_{1',2'}} = 5.7 Hz and *J*_{1',2''} = 8.3 Hz), 5.7-5.6 (m, 2H, *H*-3' and *H*-5), 4.8-4.6 (m, 3H, *H*-5', *H*-5'' and *H*-4'), 2.8-2.7 (m, 1H, *H*-2'), 2.4-2.3 (m, 1H, *H*-2'').

Reagents:

Tri-*n*-butyltin hydride: Fluka, >98%, ref 90915

Azabisisobutyronitrile: Fluka, >98%, ref 11630

3',5'-Di-O-benzoyl-2'-deoxy- β -L-4-thio-uridine (6)

A solution of compound **5** (69 g, 0.158 mol) and Lawesson's reagent (74 g) in anhydrous methylene chloride (3900 ml) was refluxed under argon overnight. After

5 evaporation of the solvent, the crude residue was purified by a silica gel column chromatography [eluant: gradient of methanol (0-2%) in methylene chloride] to afford pure compound **6** (73 g) in quantitative yield; ¹H NMR (CDCl₃) δ ppm 9.5 (br s, 1H, NH), 8.1-7.4 (m, 10H, C₆H₅CO), 7.32 (d, 1H, H-6, J = 7.7 Hz), 6.30 (dd, 1H, H-1', J = 5.6 Hz and J = 8.2 Hz), 6.22 (d, 1H, H-5, J = 7.7 Hz), 5.6 (m, 1H, H-3'), 4.7 (m, 2H, H-5', H-5''), 4.5 (m, 1H, H-4'), 2.8 (m, 1H, H-2'), 2.3 (m, 1H, H-2'').

Reagent:

15 Lawesson's reagent: Fluka, >98%, ref 61750

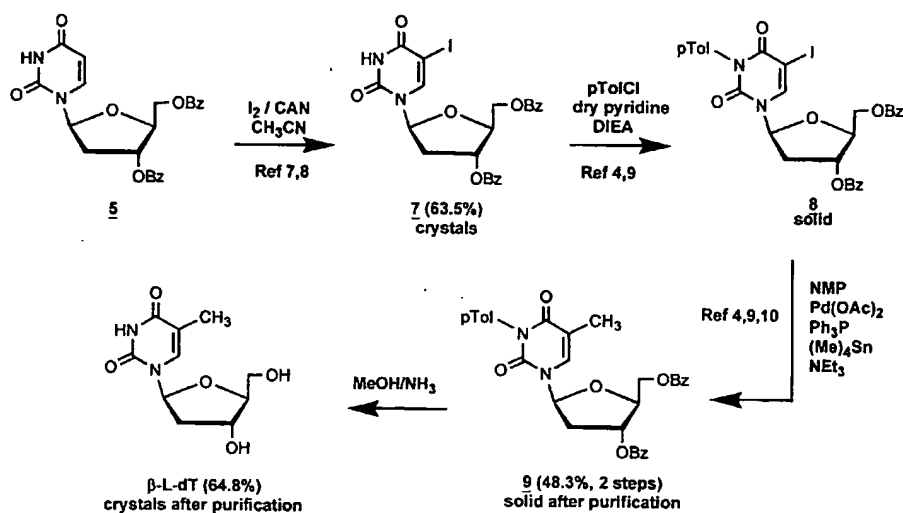
2'-Deoxy-β-L-cytosine

20 A solution of compound **6** (7.3 g, 0.016 mol) in methanol saturated with ammonia (73 ml) was heated at 100°C in a stainless steel cylinder for 3h. After cooling carefully, the solvent was evaporated under reduced pressure. An aqueous solution of the residue was washed with ethyl acetate and evaporated to dryness. Such a procedure was carried out on 9 other samples (each 7.3 g) of compound **6** (total amount of **6** = 73 g). The 10 residues were combined, diluted with absolute ethanol and cooled to give **7** as crystals. Trace of benzamide were eliminated from the crystals of **6** by a solid-liquid extraction procedure (at reflux in ethyl acetate for 1h). Yield, 28.75 g (78.6%) of compound **6**; m. p. 141-145°C; ¹H NMR (DMSO) δ ppm 8.22 and 8.00 (2 br s, 2H, NH₂), 7.98 (d, 1H, H-6, J = 7.59 Hz), 6.12 (t, 1H, H-1', J = 6.5 Hz and J = 7.6 Hz), 5.89 (d, 1H, H-5, J = 7.59 Hz), 5.3 (br s, 1H, OH-3'), 5.1 (br s, 1H, OH-5'), 4.2 (m, 1H, H-3'), 3.80 (q, 1H, H-4', J = 3.6 Hz and J = 6.9 Hz), 3.6-3.5 (m, 2H, H-5', H-5''), 2.2-2.0 (m, 2H, H-2', H-2''); FAB<0, (GT) m/e 226 (M-H)⁻, 110 (B)⁻; FAB>0 (GT) 228 (M+H)⁺, 112 (B+2H)⁺; [α]_D²⁰ - 56.48 (c = 1.08 in DMSO); UV (pH 7) λ_{max} = 270 nm (ε = 10000).

Reagent:

40 Methanolic ammonia: previously saturated at -5°C, tightly stoppered, and kept in a freezer.

Example 5 Stereoselective Synthesis of 2'-Deoxy-β-L-Thymidine (β-L-dT)



3',5'-Di-*O*-benzoyl-2'-deoxy-5-iodo-β-L-uridine (**7**)

A mixture of compound **5** (105.8 g, 0.242 mol), iodine (76.8 g), CAN (66.4 g) and acetonitrile (2550 ml) was stirred at 80°C for 3h then the reaction mixture was cooled at room temperature leading to crystallization of compound **7** (86.6 g, 63.5%); m. p. 192-194°C; ¹H NMR (DMSO) δ ppm .8.34 (s, 1H, NH), 8.2-7.2 (m, 11H, 2 C₆H₅CO, H-6), 6.31 (q, 1H, H-1', J = 5.5 Hz and J = 8.7 Hz), 5.5 (m, 1H, H-3'), 4.7 (m, 2H, H-5', H-5''), 4.5 (m, 1H, H-4'), 2.7 (m, 1H, H-2'), 2.3 (m, 1H, H-2''); FAB<0, (GT) m/e 561 (M-H)⁺, 237 (B)⁺; FAB>0 (GT) 563 (M+H)⁺; [α]_D²⁰ + 39.05 (c = 1.05 in DMSO); UV (EtOH 95) ν_{max} = 281 nm (ε = 9000), ν_{min} = 254 nm (ε = 4000), ν_{max} = 229 nm (ε = 31000); Anal. Calcd for C₂₃H₁₉IN₂O₇: C, 49.13 H, 3.41 N, 4.98 I, 22.57. Found: C, 49.31 H, 3.53 N, 5.05 I, 22.36.

Reagents:

Iodine: Fluka, 99.8%, ref 57650

Cerium ammonium nitrate (CAN): Aldrich, >98.5%, ref 21,547-3

3',5'-Di-*O*-benzoyl-2'-deoxy-3-*N*-toluoyl- β -L-thymidine (9)

To a solution of compound **7** (86.6g, 0.154 mol) in anhydrous pyridine (1530 ml) containing *N*-ethyldiisopropylamine (53.6 ml) was added, portionwise at 0°C, *p*-toluoyl chloride (40.6 ml). The reaction mixture was stirred for 2 h at room temperature, then water was added to stop the reaction and the reaction mixture was extracted with methylene chloride. The organic phase was washed with water, dried over sodium sulfate and evaporated to dryness to give crude 3',5'-di-*O*-benzoyl-2'-deoxy-3-*N*-toluoyl-5-iodo- β -L-uridine (**8**) which can be used for the next step without further purification.

A solution of the crude mixture **8**, palladium acetate (3.44 g), triphenylphosphine (8.0 g) in *N*-methylpyrrolidinone (1375 ml) with triethylamine (4.3 ml) was stirred at room temperature for 45 min. Then, tetramethyltin (42.4 ml) was added dropwise at 0°C under argon. After stirring at 100-110°C overnight, the reaction mixture was poured into water and extracted with diethyl ether. The organic solution was dried over sodium sulfate and concentrated under reduced pressure. The residue was purified by a silica gel column chromatography [eluant: stepwise gradient of ethyl acetate (0-10%) in toluene] to give compound **9** as a foam (42.3 g, 48.3% for the 2 steps). ¹H NMR (DMSO) δ ppm .8.3-7.2 (m, 15H, 2 C₆H₅CO, 1 CH₃C₆H₄CO, H-6), 6.29 (t, 1H, *H*-1', *J* = 7.0 Hz), 5.7 (m, 1H, *H*-3'), 4.7-4.5 (m, 3H, *H*-5', *H*-5'', *H*-4'), 2.7-2.6 (m, 2H, *H*-2', *H*-2''); FAB<0, (GT) *m/e* 567 (M-H)⁻, 449 (M-CH₃C₆H₄CO)⁻, 243 (B)⁻, 121 (C₆H₅COO)⁻; FAB>0 (GT) 1137 (2M+H)⁺, 569 (M+H)⁺, 325 (M-B)⁻, 245 (B+2H)⁺, 119 (CH₃C₆H₅CO)⁺.

Reagents:

***p*-Toluoyl chloride**, Aldrich, 98%, ref 10,663-1

Diisopropylethylamine: Aldrich, >99.5%, ref 38,764-9

***N*-methylpyrrolidinone**: Aldrich, >99%, ref 44,377-8

Palladium acetate: Aldrich, >99.98%, ref 37,987-5

Triphenylphosphine: Fluka, >97%, ref 93092

Tetramethyltin: Aldrich, >99%, ref 14,647-1

2'-Deoxy- β -L-thymidine

A solution of compound **9** (42.3 g, 0.074 mol) in methanol saturated with ammonia (1850 ml) was stirred at room temperature for two days. After evaporation of the solvent, the residue was diluted with water and washed several times with ethyl acetate. The aqueous layer was

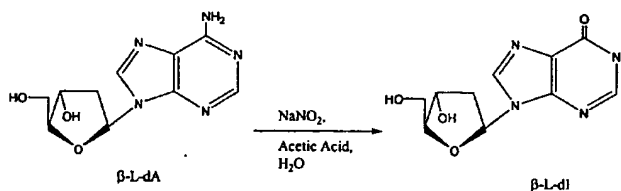
separated, evaporated under reduced pressure and the residue was purified by a silica gel column chromatography [eluant: stepwise gradient of methanol (0-10%) in methylene chloride] to give pure 2'-deoxy- β -L-thymidine (11.62 g, 64.8%) which was crystallized from ethanol; m.p. 185-188°C; ^1H NMR (DMSO) δ ppm 11.3 (s, 1H, NH), 7.70 (s, 1H, H-6), 6.2 (pt, 1H, H-1'), 5.24 (d, 1H, OH-3', J = 4.2 Hz), 5.08 (t, 1H, OH-5', J = 5.1 Hz), 4.2 (m, 1H, H-3'), 3.7 (m, 1H, H-4'), 3.5-3.6 (m, 2H, H-5', H-5''), 2.1-2.0 (m, 2H, H-2', H-2''); FAB<0, (GT) m/e 483 (2M-H)⁻, 349 (M+T-H)⁻, 241 (M-H)⁻, 125 (B)⁻; FAB>0 (GT) 243 (M+H)⁺, 127 (B+2H)⁺;)⁺; [α]_D²⁰ - 13.0 (c = 1.0 in DMSO); UV (pH 1) ν_{max} = 267 nm (ϵ = 9700), ν_{min} = 234 nm (ϵ = 2000).

Reagent:

Methanolic ammonia: previously saturated at -5°C, tightly stoppered, and kept in a freezer.

Example 6 Stereoselective Synthesis of 2'-deoxy- β -L-inosine (β -L-dI)

β -L-dI was synthesized by deamination of 2'-deoxy- β -L-adenosine (β -L-dA) following a procedure previously described in the 9-D-glucopyranosyl series (Ref: I. Iwai, T. Nishimura and B. Shimizu, Synthetic Procedures in Nucleic Acid Chemistry, W. W. Aorbach and R. S. Tipson, eds., John Wiley & Sons, Inc. New York, vol. 1, pp. 135-138 (1968)).



Thus, a solution of β -L-dA (200 mg) in a mixture of acetic acid (0.61 ml) and water (19 ml) was heated with sodium nitrite (495 mg), and the mixture was stirred at room temperature overnight. The solution was then evaporated to dryness under diminished pressure. An aqueous solution of the residue was applied to a column of IR-120 (H^+) ion-exchange resin, and the column was eluted with water. Appropriate fractions were collected and evaporated to dryness to afford pure β -L-dI which was crystallized from methanol (106 mg, 53% yield not optimized): m.p.=209°-211°C; UV (H_2O), λ_{max} =247 nm; ^1H -NMR

(DMSO-d₆)= 8.32 and 8.07 (2s, 1H each, H-2 and H-8), 6.32 (pt, 1H, H-1; J=6.7 Hz), 4.4 (m, 1H, H-3'), 3.9 (m, 1H, H-4'), 3.7-3.4 (m, 2H partially obscured by HOD, H-5',5''), 2.6 and 2.3 (2m, 1H each, H-2' and H-2''); mass spectra (mature, glycerol-thioglycerol, 1:1, v/v), FAB>0: 253 (m+H)⁺, 137 (base + 2H)⁺; FAB<0: 251 (m-H)⁻, 135 (base)⁻; [α]_D²⁰= +19.3 (-c 0.88, H₂O).

Anti-HBV Activity of the Active Compounds

The ability of the active compounds to inhibit the growth of virus in 2.2.15 cell cultures (HepG2 cells transformed with hepatitis virion) can be evaluated as described in detail below.

A summary and description of the assay for antiviral effects in this culture system and the analysis of HBV DNA has been described (Korba and Milman, 1991, Antiviral Res., 15:217). The antiviral evaluations are performed on two separate passages of cells. All wells, in all plates, are seeded at the same density and at the same time.

Due to the inherent variations in the levels of both intracellular and extracellular HBV DNA, only depressions greater than 3.5-fold (for HBV virion DNA) or 3.0-fold (for HBV DNA replication intermediates) from the average levels for these HBV DNA forms in untreated cells are considered to be statistically significant (P<0.05). The levels of integrated HBV DNA in each cellular DNA preparation (which remain constant on a per cell basis in these experiments) are used to calculate the levels of intracellular HBV DNA forms, thereby ensuring that equal amounts of cellular DNA are compared between separate samples.

Typical values for extracellular HBV virion DNA in untreated cells range from 50 to 150 pg/ml culture medium (average of approximately 76 pg/ml). Intracellular HBV DNA replication intermediates in untreated cells range from 50 to 100 μg/pg cell DNA (average approximately 74 pg/μg cell DNA). In general, depressions in the levels of intracellular HBV DNA due to treatment with antiviral compounds are less pronounced, and occur more slowly, than depressions in the levels of HBV virion DNA (Korba and Milman, 1991, Antiviral Res., 15:217).

The manner in which the hybridization analyses are performed for these experiments result in an equivalence of approximately 1.0 pg of intracellular HBV DNA to 2-3 genomic copies per cell and 1.0 pg/ml of extracellular HBV DNA to 3 x 10⁵ viral particles/ml.

Example 7

The ability of the triphosphate derivatives of β -L-dA, β -L-dC, β -L-dU, β -L-2'-dG, β -L-dI, and β -L-dT to inhibit hepatitis B was tested. Table 1 describes the comparative inhibitory activities of triphosphates of β -L-dT (β -L-dT-TP), β -L-dC (β -L-dC-TP), β -L-dU (β -L-dU-TP) and β -L-dA (β -L-dA-TP) on woodchuck hepatitis virus (WHV) DNA polymerase, human DNA polymerases α , β , and γ .

Table 1

Inhibitor	WHV DNA pol IC ₅₀	DNA pol α K _i ^b (μ M)	DNA pol β K _i ^b (μ M)	DNA pol γ K _i ^b (μ M)
β -L-dT-TP	0.34	>100	>100	>100
β -L-dA-TP	2.3	>100	>100	>100
β -L-dC-TP	2.0	>100	>100	>100
β -L-dU-TP	8	>100	>100	>100

^aIC₅₀: 50% Inhibitory concentration

^bK_i value was determined using calf thymus activated DNA as template-primer and dATP as substrate. Inhibitors were analyzed by Dixon plot analysis. Under these conditions, the calculated mean K_m of human DNA polymerase α for dATP as approximately 2.6 μ M. Human DNA polymerase β exhibited a steady state K_m of 3.33 μ M for dATP. Human DNA polymerase γ exhibited a steady K_m of 5.2 μ M.

Example 8

The anti-hepatitis B virus activity of β -L-dA, β -L-dC, β -L-dU, β -L-2'-dG and β -L-dT was tested in transfected Hep G-2 (2.2.15) cells. Table 2 illustrates the effect of β -L-dA, β -L-dC, β -L-dU, and β -L-dT against hepatitis B virus replication in transfected Hep G-2 (2.2.15) cells.

Table 2

Compound	HBV virions ^a EC ₅₀ (μ M)	HBV Ri ^b EC ₅₀ (μ M)	Cytotoxicity IC ₅₀ (μ M)	Selectivity Index IC ₅₀ /EC ₅₀
β -L-dT	0.05	0.05	>200	>4000
β -L-dC	0.05	0.05	>200	>4000
β -L-dA	0.10	0.10	>200	>2000

Compound	HBV virions ^a EC ₅₀ (μM)	HBV Ri ^b EC ₅₀ (μM)	Cytotoxicity IC ₅₀ (μM)	Selectivity Index IC ₅₀ /EC ₅₀
β-L-dI	1.0	1.0	>200	>200
β-L-dU	5.0	5.0	>200	>40

^aExtracellular DNA^bReplicative intermediates (Intracellular DNA)**Example 9**

The effect of β-L-dA, β-L-dC and β-L-dT in combination on the growth of hepatitis B was measured in 2.2.15 cells. The results are provided in Table 3.

Table 3

Combination	Ratio	EC ₅₀
L-dC + L-dT	1:3	.023
L-dC + L-dT	1:1	.053
L-dC + L-dT	3:1	.039
L-dC + L-dA	1:30	.022
L-dC + L-dA	1:10	.041
L-dC + L-dA	1:3	.075
L-dT + L-dA	1:30	.054
L-dT + L-dA	1:10	.077
L-dT + L-dA	1:3	.035

Each combination produced anti-HBV activity that was synergistic. In addition, the combination of L-dA + L-dC + L-dT was also synergistic in this model.

Example 10

The inhibition of hepatitis B replication in 2.2.15 cells by β-L-dA and β-L-dC, alone and in combination was measured. The results are shown in Table 4.

Table 4

^a β-L-2'-deoxy-adenosine (μM)	^b β-L-2'-deoxy-cytidine (μM)	% Inhibition	^c C.I.
0.5		90	
0.05		24	
0.005		1	
	0.5	95	
	0.05	40	
	0.005	10	
0.05	0.05	80	0.34
0.05	0.005	56	0.20
0.05	0.0005	50	0.56
0.005	0.05	72	0.35
0.005	0.005	54	0.35
0.005	0.0005	30	0.16
0.0005	0.05	50	0.83
0.0005	0.005	15	0.28
0.0005	0.0005	0	N.A.

^aβ-L-2'-deoxy-adenosine: IC₅₀=0.09 μM

^bβ-L-2'-deoxy-cytidine: IC₅₀=0.06 μM

^cCombination indices values indicate synergism effect (<1), additive effect (=1), and antagonism effect (>1)

Example 11

The efficacy of L-dA, L-dT and L-dC against hepadnavirus infection in woodchucks (*Marmota monax*) chronically infected with woodchuck hepatitis virus (WHV) was determined. This animal model of HBV infection is widely accepted and has proven to be useful for the evaluation of antiviral agents directed against HBV.

Protocol:

Experimental groups (n=3 animals/drug group, n=4 animals/control)

Group 1	vehicle control
Group 2	lamivudine (3TC) (10 mg/kg/day)
Groups 3-6	L-dA (0.01, 0.1, 1.0, 10 mg/kg/day)
Groups 7-10	L-dT (0.01, 0.1, 1.0, 10 mg/kg/day)
Groups 11-14	L-dC (0.01, 0.1, 1.0, 10 mg/kg/day)

Drugs were administered by oral gavage once daily, and blood samples taken on days 0, 1, 3, 7, 14, 21, 28, and on post-treatment days +1, +3, +7, +14, +28 and +56. Assessment

of the activity and toxicity was based on the reduction of WHV DNA in serum: dot-blot, quantitative PCR.

The results are illustrated in Figure 3 and Table 5.

Table 5
Antiviral Activity of LdA, LdT, LdC in Woodchuck Model of Chronic HBV Infection

day	Control	LdA	LdT	LdC
	ng WHV-DNA per ml serum ^{1,2}			
0	381	436	423	426
1	398	369	45	123
3	412	140	14	62
7	446	102	6	46
14	392	74	1	20

¹ LdA, LdT, LdC administered orally once a day at 10mg/kg

² limit of detection is 1 ng/ml WHV-DNA per ml serum

The data show that L-dA, L-dT and L-dC are highly active in this in vivo model. First, viral load is reduced to undetectable (L-dT) or nearly undetectable (L-dA, L-dC) levels. Second, L-dA, L-dT and L-dC are shown to be more active than 3TC (lamivudine) in this model. Third, viral rebound is not detected for at least two weeks after withdrawal of L-dT. Fourth, dose response curves suggest that a modes increase in the dose of L-dA and L-dC would show antiviral activity similar to L-dT. Fifth, all animals receiving the drugs gained weight and no drug-related toxicity was detected.

Toxicity of Compounds

Toxicity analyses were performed to assess whether any observed antiviral effects are due to a general effect on cell viability. The method used is the measurement of the effect of β -L-dA, β -L-dC and β -L-dT on cell growth in human bone marrow clorogenic assays, as compared to Lamuvudine. The results are provided in Table 6.

Table 6

Compound	CFU-GM (μ M)	BFU-E (μ M)
β -L-dA	>10	>10
β -L-dC	>10	>10
β -L-dT	>10	>10
β -L-dU	>10	>10
Lamuvudine	>10	>10

Preparation of Pharmaceutical Compositions

Humans suffering from any of the disorders described herein, including hepatitis B, can be treated by administering to the patient an effective amount of a β -2'-deoxy- β -L-erythro-pentofuranonucleoside, for example, β -L-2'-deoxyadenosine, β -L-2'-deoxycytidine, β -L-2'-deoxyuridine, β -L-2'-deoxyguanosine or β -L-2'-deoxythymidine or a pharmaceutically acceptable prodrug or salt thereof in the presence of a pharmaceutically acceptable carrier or diluent. The active materials can be administered by any appropriate route, for example, orally, parenterally, intravenously, intradermally, subcutaneously, or topically, in liquid or solid form.

The active compound is included in the pharmaceutically acceptable carrier or diluent in an amount sufficient to deliver to a patient a therapeutically effective amount of compound to inhibit viral replication *in vivo*, without causing serious toxic effects in the patient treated. By "inhibitory amount" is meant an amount of active ingredient sufficient to exert an inhibitory effect as measured by, for example, an assay such as the ones described herein.

A preferred dose of the compound for all of the abovementioned conditions will be in the range from about 1 to 50 mg/kg, preferably 1 to 20 mg/kg, of body weight per day, more generally 0.1 to about 100 mg per kilogram body weight of the recipient per day. The effective dosage range of the pharmaceutically acceptable prodrug can be calculated based on the weight of the parent nucleoside to be delivered. If the prodrug exhibits activity in itself, the effective dosage can be estimated as above using the weight of the prodrug, or by other means known to those skilled in the art.

5 The compound is conveniently administered in unit any suitable dosage form, including but not limited to one containing 7 to 3000 mg, preferably 70 to 1400 mg of active ingredient per unit dosage form. A oral dosage of 50-1000 mg is usually convenient.

10 Ideally the active ingredient should be administered to achieve peak plasma concentrations of the active compound of from about 0.2 to 70 μ M, preferably about 1.0 to 10 μ M. This may be achieved, for example, by the intravenous injection of a 0.1 to 5% solution of the active ingredient, optionally in saline, or administered as a bolus of the active ingredient.

15 The concentration of active compound in the drug composition will depend on absorption, inactivation, and excretion rates of the drug as well as other factors known to those of skill in the art. It is to be noted that dosage values will also vary with the severity of the condition to be alleviated. It is to be further understood that for any particular subject, specific dosage regimens should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the compositions, and that the concentration ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the claimed composition. The active ingredient may be administered at once, or may be divided into a number of smaller doses to be administered at varying intervals of time.

30 A preferred mode of administration of the active compound is oral. Oral compositions will generally include an inert diluent or an edible carrier. They may be enclosed in gelatin capsules or compressed into tablets. For the purpose of oral therapeutic administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition.

40 The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring. When the dosage unit form is a capsule, it can contain, in addition to material of the above type, a liquid carrier such as a fatty oil. In addition, dosage unit forms can contain various other materials which

5 modify the physical form of the dosage unit, for example, coatings of sugar, shellac, or other enteric agents.

The compound can be administered as a component of an elixir, suspension, syrup, wafer, chewing gum or the like. A syrup may contain, in addition to the active compounds, sucrose as a sweetening agent and certain preservatives, dyes and colorings and flavors.

10 The compound or a pharmaceutically acceptable derivative or salts thereof can also be mixed with other active materials that do not impair the desired action, or with materials that supplement the desired action, such as antibiotics, antifungals, antiinflammatories, protease inhibitors, or other nucleoside or nonnucleoside antiviral agents. Solutions or suspensions used for parenteral, intradermal, subcutaneous, or topical application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid; buffers such as acetates, citrates or phosphates and agents for the adjustment of tonicity such as sodium chloride or dextrose. The parental preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

25 If administered intravenously, preferred carriers are physiological saline or phosphate buffered saline (PBS).

30 In a preferred embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation.

40 Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) are also preferred as pharmaceutically acceptable carriers. These may be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811. For example, liposome formulations may be prepared by dissolving appropriate lipid(s) (such as stearyl phosphatidyl ethanolamine, stearyl phosphatidyl choline, arachadoyl phosphatidyl choline, and

5 cholesterol) in an inorganic solvent that is then evaporated, leaving behind a thin film of dried
lipid on the surface of the container. An aqueous solution of the active compound or its
monophosphate, diphosphate, and/or triphosphate derivatives is then introduced into the
10 container. The container is then swirled by hand to free lipid material from the sides of the
container and to disperse lipid aggregates, thereby forming the liposomal suspension.

This invention has been described with reference to its preferred embodiments.
Variations and modifications of the invention, will be obvious to those skilled in the art from
15 the foregoing detailed description of the invention. It is intended that all of these variations
and modifications be included within the scope of the this invention.

Claims

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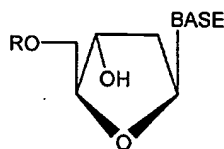
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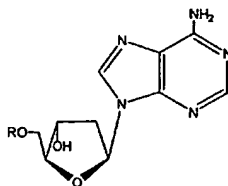
We claim:

1. A method for treating a host infected with hepatitis B comprising administering an effective amount of an anti-HBV biologically active 2'-deoxy- β -L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside has the formula:



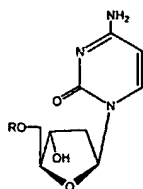
wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may be optionally substituted.

2. The method of claim 1, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyadenosine or a pharmaceutically acceptable salt or prodrug thereof, of the formula:



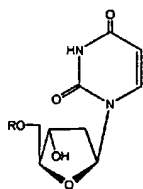
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

3. The method of claim 1, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxycytidine or pharmaceutically acceptable salt or prodrug thereof of the formula:



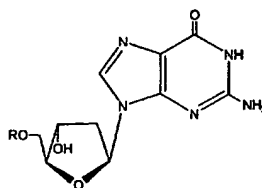
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

4. The method of claim 1, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyuridine or pharmaceutically acceptable salt or prodrug thereof of the formula:



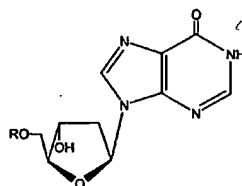
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

5. The method of claim 1, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyguanosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



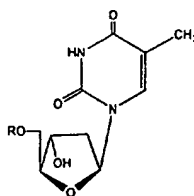
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

6. The method of claim 1, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyinosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



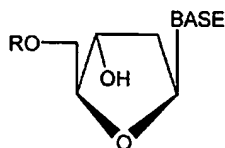
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

7. The method of claim 1, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-thymidine or a pharmaceutically acceptable salt or prodrug thereof of the formula:



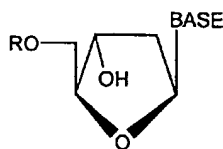
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

8. A method for treating a host infected with hepatitis B comprising administering an effective amount of two or more anti-HBV biologically active 2'-deoxy- β -L-erythro-pentofuranonucleosides or a pharmaceutically acceptable salt or prodrug thereof in combination of alternation, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleosides have the formula:



wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may be optionally substituted.

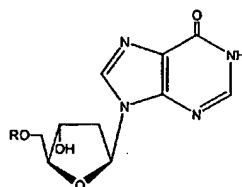
9. A method for treating a host infected with hepatitis B comprising administering an effective amount of an anti-HBV biologically active 2'-deoxy-β-L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof in combination or alternation with an additional anti-hepatitis B agent, wherein the 2'-deoxy-β-L-erythro-pentofuranonucleoside has the formula:



wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may be optionally substituted.

10. The method of claim 9, wherein the additional anti-hepatitis B agent is selected from the group consisting of 3TC, FTC, L-FMAU, DAPD, famciclovir, penciclovir, BMS-200475, bis pom PMEA (adefovir, dipivoxil), lobucavir, ganciclovir, or ribavarin

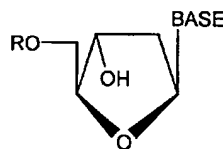
11. A compound or pharmaceutically acceptable salt or prodrug thereof of the formula:



wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

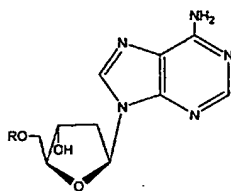
12. A pharmaceutical composition comprising an effective amount of a compound of claim 10 in combination with a pharmaceutically acceptable carrier.

13. Use of an anti-HBV biologically active 2'-deoxy- β -L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof for the treatment of a host infected with hepatitis B, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside has the formula:



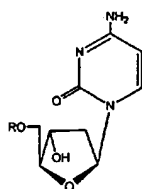
wherein R is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, CO-aryloxyalkyl, CO-substituted aryl, alkylsulfonyl, arylsulfonyl, aralkylsulfonyl, amino acid residue, mono, di, or triphosphate, or a phosphate derivative; and BASE is a purine or pyrimidine base which may be optionally substituted.

14. The use of claim 13, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyadenosine or a pharmaceutically acceptable salt or prodrug thereof, of the formula:



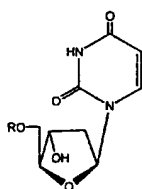
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

15. The use of claim 13, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxycytidine or pharmaceutically acceptable salt or prodrug thereof of the formula:



wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

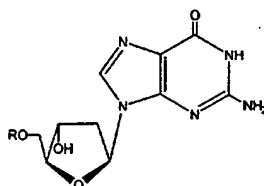
16. The use of claim 13, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside



is β -L-2'-deoxyuridine or pharmaceutically acceptable salt or prodrug thereof of the formula:

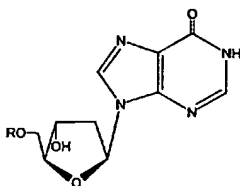
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

17. The use of claim 13, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyguanosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



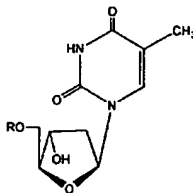
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

18. The use of claim 13, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyinosine or pharmaceutically acceptable salt or prodrug thereof of the formula:

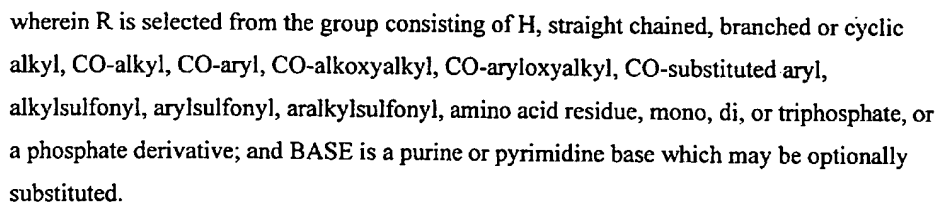


wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

19. The use of claim 13, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-thymidine or a pharmaceutically acceptable salt or prodrug thereof of the formula:

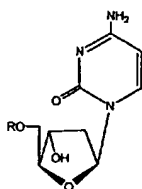


20. Use of an anti-HBV biologically active 2'-deoxy- β -L-erythro-pentofuranonucleoside or a pharmaceutically acceptable salt or prodrug thereof for the preparation of a medicament for the treatment of hepatitis B, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside has the formula:

NC1=NC=NC2=C1N=CN2C3=CN=CN=C3C4C(CO4)OC(RO)CO

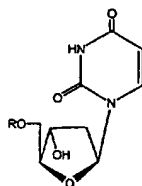
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

22. The use of claim 20, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxycytidine or pharmaceutically acceptable salt or prodrug thereof of the formula:



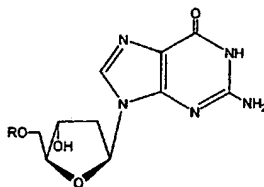
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

23. The use of claim 20, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyuridine or pharmaceutically acceptable salt or prodrug thereof of the formula:



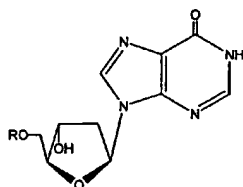
wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative (to form a stabilized nucleotide prodrug).

24. The use of claim 20, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside is β -L-2'-deoxyguanosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



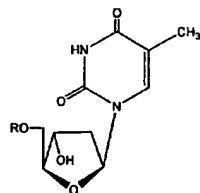
5 wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative
(to form a stabilized nucleotide prodrug).

10 25. The use of claim 20, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside
is β -L-2'-deoxyinosine or pharmaceutically acceptable salt or prodrug thereof of the formula:



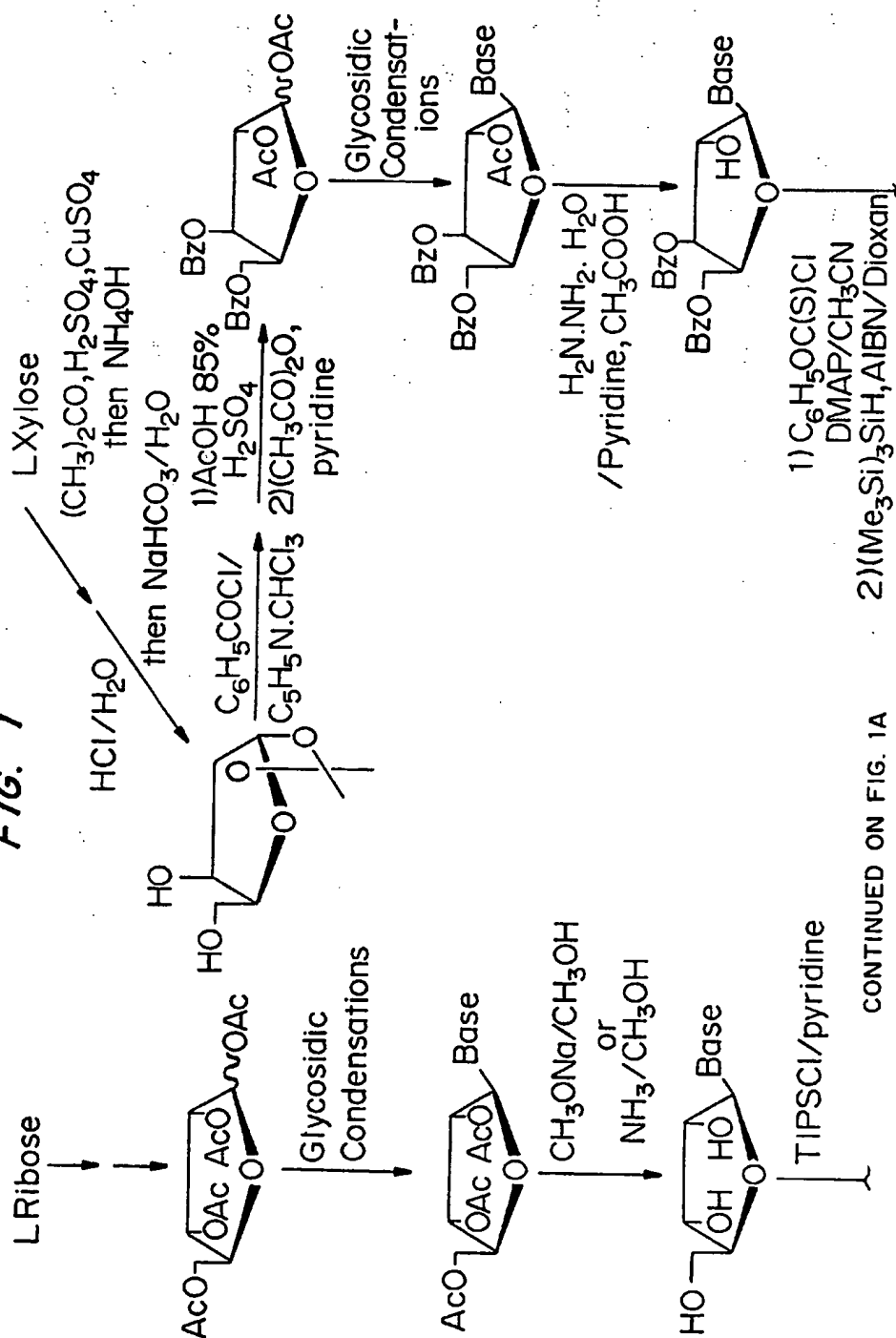
20 wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative
(to form a stabilized nucleotide prodrug).

25 26. The use of claim 20, wherein the 2'-deoxy- β -L-erythro-pentofuranonucleoside
is β -L-thymidine or a pharmaceutically acceptable salt or prodrug thereof of the formula:



35 wherein R is H, mono, di or tri phosphate, acyl, or alkyl, or a stabilized phosphate derivative
(to form a stabilized nucleotide prodrug).

FIG. 1



CONTINUED ON FIG. 1A

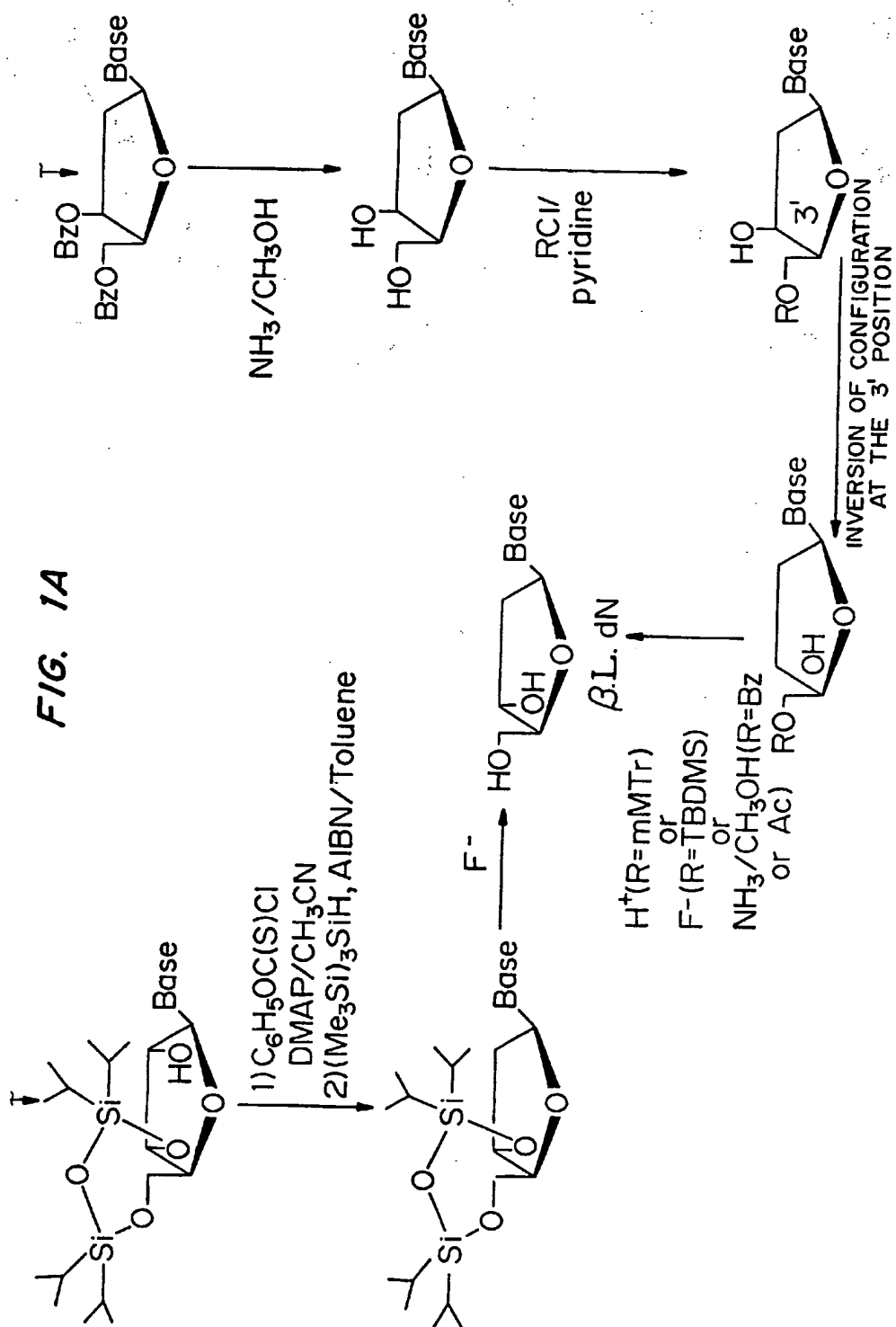
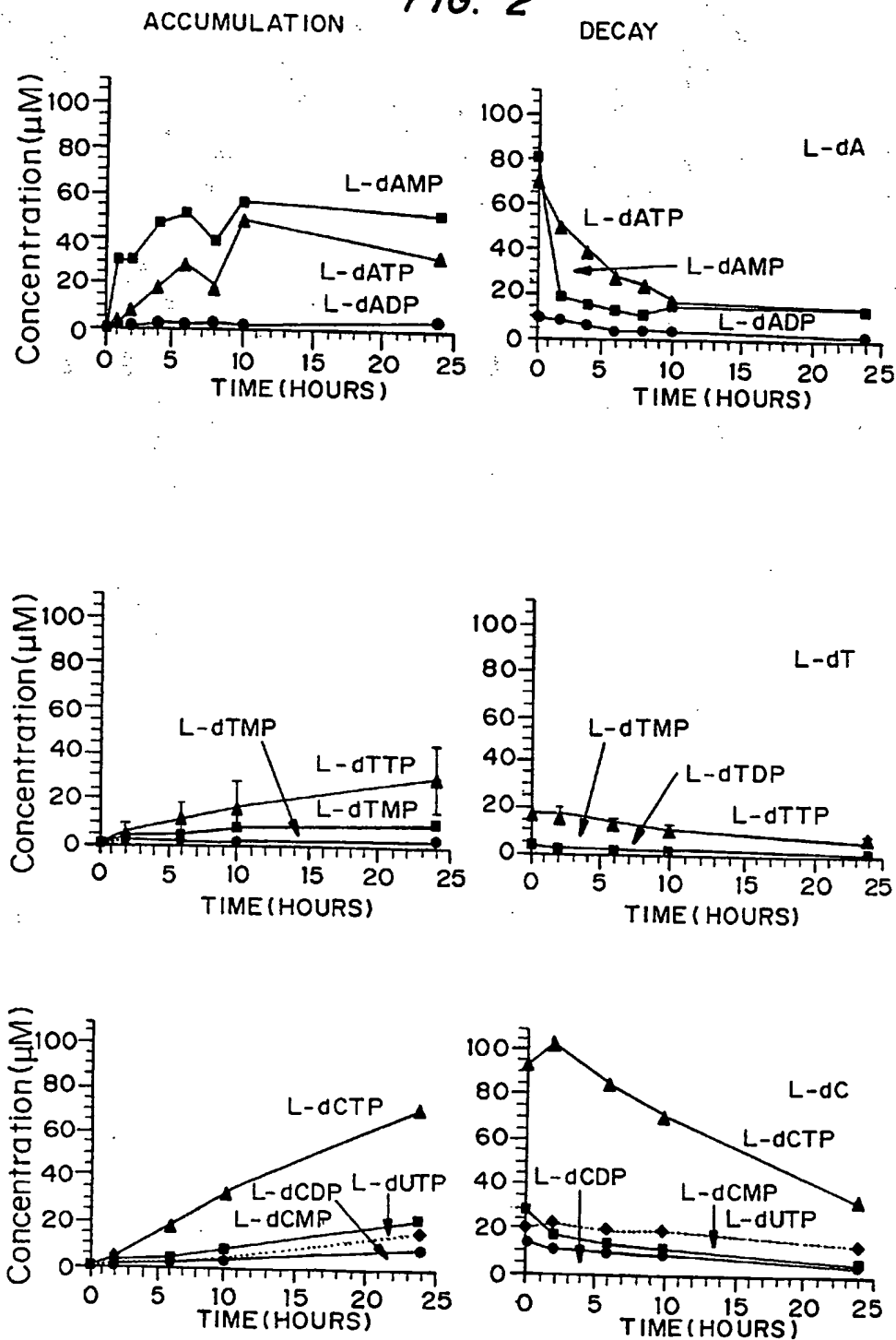


FIG. 2



(n=3 per drug treatment group, n=4 per placebo group, dose 10mg/kg orally once per day)

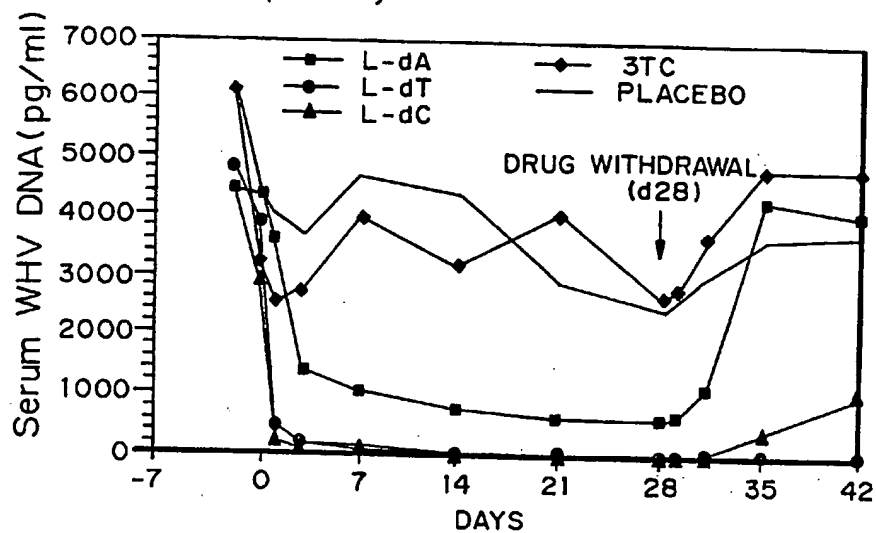


FIG. 3